

Genetic Correspondence and Comparative Spatial Analysis with Age and Gender Specification of ICMR Data on Cancer Incidence in India

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Abstract

Cancer remains a significant global health challenge, characterized by immense suffering and often limited treatment options. We aim to analyze genetic data related to five common cancer types, each presenting unique challenges and implications for diagnosis and treatment. Our goal is to identify the probable genetic causes underlying these cancers, providing leads for early identification and reducing fatality rates. Focusing on India, we explore the spatial and genetic aspects of cancer, utilizing statistical concepts and analyzing data from specific years. We aim to shed light on cancer prevalence, distribution and genetic underpinnings within the Indian population. Additionally, we investigate gender-specific and age-specific cancer incidence to offer detailed insights into the cancer landscape in India.

Dataset

In this project, we deal with multiple datasets that are based on cancer patients and the nature of cancer in both male and female human bodies.

1. Cancer causing genes dataset: The input dataset contains 802 samples corresponding to 802 people who have been detected with different types of cancer. Each sample contains expression values of more than 20K genes. Samples are categorized into one of the following types of tumors: BRCA, KIRC, COAD, LUAD and PRAD.

Here's an extended description of the tumors associated with this dataset:

- **BRCA (Breast Cancer):** Breast cancer is one of the most common cancers among women. It originates in the breast tissue and can occur in both men and women, although it is more prevalent in women. The

dataset likely includes samples from individuals with breast cancer, and the analysis aims to identify the genes associated with this type of cancer.

- KIRC (Renal Cancer): Renal cell carcinoma, or kidney cancer, occurs in the lining of small tubes in the kidney. It is one of the common forms of kidney cancer. The dataset may contain samples from individuals with renal cancer, and the analysis aims to uncover the genetic factors associated with this type of cancer.
- COAD (Colon Cancer): Colorectal cancer includes colon cancer (affecting the large intestine) and rectal cancer (affecting the rectum). Colon cancer is a major cause of cancer-related deaths. The dataset could include samples from individuals with colon cancer, with a focus on identifying the genes responsible for this type of cancer.
- LUAD (Lung Cancer - Adenocarcinoma): Lung *adenocarcinoma* is a subtype of non-small cell lung cancer. It originates in the cells lining the airways and is one of the most common types of lung cancer. The dataset may contain samples from individuals with lung adenocarcinoma and the analysis aims to uncover the genetic factors specific to this form of lung cancer.
- PRAD (Prostate Cancer): Prostate cancer occurs in the prostate, a small gland in men that produces seminal fluid. It is one of the most common cancers in men. The dataset could include samples from individuals with prostate cancer, focusing on identifying the genes associated with this type of cancer.

2. Estimated State-wise cancer incidences: The table provides estimated incidence of cancer cases in India by States and Union Territory-wise for all sites and for both sexes.

3. Gender-wise different sites of cancer: The table presents the estimated cancer incidence, number of cases, crude rate, and cumulative risk by sex and anatomical sites in India for the year 2022.

Here's an extended description of the terminologies associated with this dataset:

- Cum-risk: Cumulative risk of developing cancer in the age range of 0 to 74 years. It represents the likelihood or probability of an individual developing cancer within this specified age range.

$$\text{Cumulative Risk} = \frac{\text{Number of Cancer Cases}}{\text{Total Population}} \times 100$$

- Crude Rate (CR): Crude rate refers to the total number of cancer cases occurring in a population divided by the total population, expressed as a rate per a specific unit of time (usually per 100,000 population). It provides a general overview of cancer incidence within a population, without considering factors such as age distribution.

$$\text{Crude Rate} = \frac{\text{Number of Cancer Cases}}{\text{Total Population}} \times 100,000$$

- Age-Adjusted Rate (AAR): Age adjusted rate is a standardized rate that takes into account the age distribution of a population. It allows for fair comparisons of cancer rates between different populations or over time by removing the influence of age as a confounding factor.

$$\text{Age-Adjusted Rate} = \sum_{i=1}^n \left(\frac{\text{Age-specific rate}_i \times \text{Population}_i}{\text{Total Population}} \right) \times 100,000$$

- Malig Imm.Prol D (Malignant Immunoproliferative Diseases): This term refers to a group of disorders characterized by the abnormal proliferation of immune cells, leading to the development of malignancies. These diseases involve the uncontrolled growth of cells of the immune system, such as lymphocytes or plasma cells, and can include conditions like lymphomas, leukemias and multiple myeloma.

4. Age-wise cancer incidences: The table provides gender-disaggregated, estimated top five leading sites of cancer (%) in India by age group (0 – 14, 15 – 39, 40 – 64 & 65+ age groups) for the year 2022.

Data Pre-processing and Cleaning

We conducted data pre-processing and cleaning tasks on two datasets: “`data.csv`” and “`labels.csv`”. These datasets contain information related to cancer research, focusing specially on genetic data and associated labels.

1. Essential Packages: For this project, we utilized several essential packages in R, including `tibble`, `tidyverse` and `dplyr`.

2. Pre-processing: This step is very crucial pertaining to the complexity and volume of genetic data involved.

- Data Importing: Our code reads the CSV files “`data.csv`” and “`labels.csv`” into R as tibbles using the `read.csv()` function. These datasets contain genetic data and label information related to cancer research.
- Exploratory Data Analysis (EDA): We performed EDA to gain insights into the structure and content of the datasets. This involved examining the structure of both datasets using the `str()` function and displaying the first few entries of each dataset using the `head()` function.
- Categorical Variable Conversion: We converted some columns in the datasets from *character* type to *factor* type using the `as.factor()` function. This was necessary for categorical variables. We also converted the final dataframe into a more structured format, *tibble* to facilitate efficient analysis.
- Data Merging: We merged the two datasets into a single dataframe using the `cbind()` function, combining the label information with the genetic data.
- Data Quality Checks: We checked for null values (`is.null()`), NA values (`is.na()`), and duplicated entries (`duplicated()`) in the merged dataframe. The absence of such issues indicates clean data.
- Summary Statistics: We calculated summary statistics for the genetic data to understand its distribution and characteristics using the `summary()` function.

The processed dataframe contains both label information and genetic data. It is saved in the “`.Rdata`” format for further analysis and modelling.

Statistical Data Analysis:

The data we have, has 20,534 columns with 800 rows, as mentioned in the description. And distribution of classes is given below:

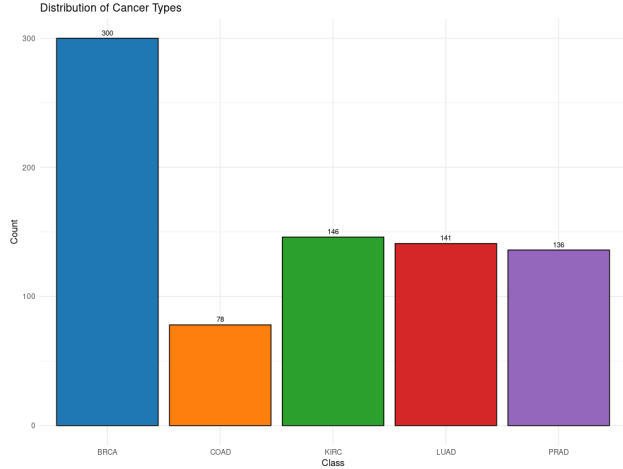


Figure 1: Frequency of Classes

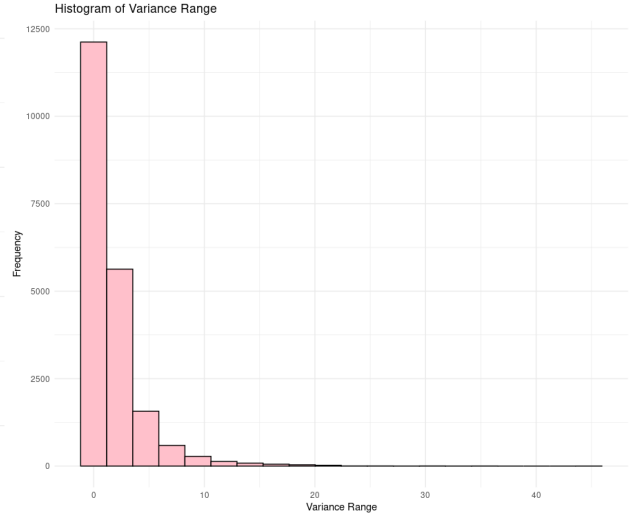


Figure 2: Histogram of Classes

From the frequency plot, we can see that instances for class BRCA are more than that of others followed by KIRC, LUAD and PRAD and class COAD has the least number of instances.

- Variance Threshold: From the graph, it's easy to see that most of the columns have low variance. Features with high variance have data points that are spread out over a wide range. These features are considered significant because they contain valuable information that can help in discriminating between different classes or categories. Similarly, features with low variance have data points that have little variation. Such features may not carry much discriminative power and can be considered less significant.
- Age specific genderwise cancer incidences: The shapes of the density curves represents the leptokurtic curve for females and mesokurtic curve for males.

