# Sequence API Assignment

#### Overview

Genetic variant classification systems predict the pathological consequences of a particular sequence variant (i.e. change in one, or several, basepairs of a genome) by comparing genomic changes caused by that variant to a reference genome. The goal of this assignment is to build an API which will provide programmatic access to the reference Human genome.

A copy of the reference genome (build GRCh37) is available here: https://s3.amazonaws.com/downloads.solvebio.com/sequence/genbank.GRCh 37.fa.gz

Please note that the .fa extension denotes FastA file format. FastA is a very common file format for storing genomes; please feel free to use any third-party Python FastA libraries that you can find.

## **API Design**

Your API should return a portion of the Human genome specified by an given genetic coordinate range. Genomic ranges are composed of chromosome, start and stop, where positions are given relative to the first basepair of a chromosome. **NOTE: genomic coordinates start at 1.** 

Detailed specifications are given below.

#### Request

Method	Parameters
GET	<pre>{     chromosome: <str>,     start: <integer>,     stop: <integer>}</integer></integer></str></pre>
POST	<pre>chromosome: <str>,     start: <integer>,     stop: <integer> }, </integer></integer></str></pre>

 ${\tt POST}$  should allow a user to retrieve multiple sequence ranges in the same request.

Parameter	Python Type	Value
chromosome	str	1-22, X, Y
start	int	[1,]
stop	int	[1,]

#### Response

Status Code	Body
200	a, or many (if POST), genetic sequence(s) (str containing A, T, G, C's)
400	detailed error response

## Requirements

Your API should be built on Django and Django REST Framework, it should be REST-ful, should return JSON documents, and should conform to standard conventions and best-practices. Additionally, it should **only** return HTTP 200 and HTTP 400 status codes. All other responses should be handled server-side.

#### Additionally:

- the API should be one-based, fully-closed (i.e. it should support inclusive ranges)
- the API should support request ranges of up 500 basepairs; requests for longer ranges should result in an HTTP 400
- the API should not support negative ranges (i.e. [10, 8], ranges where start > stop)

### **Submission and Deliverables:**

Please provide a link to your GitHub repository with the API, a README with complete documentation, and corresponding tests.

Bonus: Add support for additional genome builds (versions).