

HUMAN GENETIC VARIATION VIEWER

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- Motivation
- Solution
- Demo and Use-Cases
- Implementation
- Improvements

MOTIVATION

The power of the unaided mind is highly overrated. The real powers come from devising external aids that enhance cognitive abilities. – Donald Norman

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- Exploratory visualization is the first step towards discovering patterns
- Variation viewers are *absent*, if not, provide limited flexibility

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- Scalable, and Interactive exploration on the browser

DEMO

DETAILS

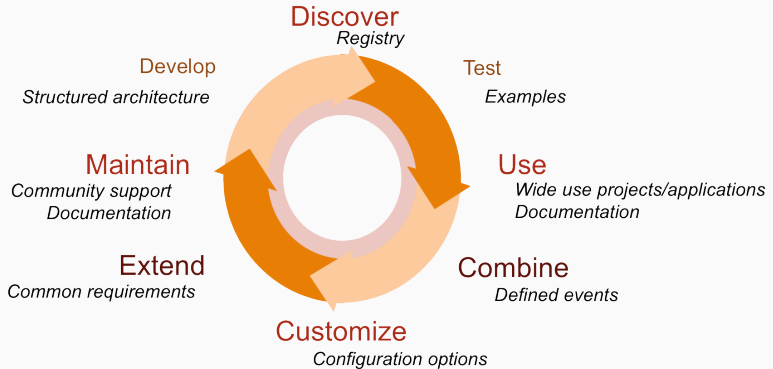
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- Events system that triggers events on user actions, allows cross-component communication

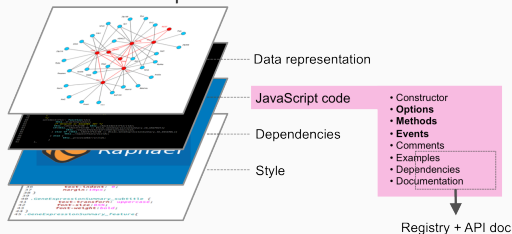
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WHY BIOJS

Reusable components that can talk to each other



```
{  
  "id": "P00533_variant226",  
  "sourceIds": ["COSM1090877", "COSM1090879"],  
  "position": 541,  
  "wild_type": "L",  
  "mutation": "I",  
  "frequency": 0.0,  
  "polyphenPrediction": "benign",  
  "polyphenScore": 0.0,  
  "siftPrediction": "tolerated",  
  "siftScore": 0.86,  
  "somaticStatus": 1,  
  "consequenceTypes": "missense variant",  
  "cytogeneticBand": "7p11.2",  
  "genomicLocation": "7:g.55229314C>A"  
}
```

- Pre-generated JSON files
- Current version uses files generated by an unpublished webservice at EBI
- Protein variants, though not specific to it

- Supports JSON formatted files, alpha VCF support
- User defined scoring criteria
- Different levels of information
 - Overview: Condensed information
 - Zoomed View: All annotations
- Loading proteins through URL parameters
- SIFT, Polyphen,

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- Benchmarking predictions

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- Interaction with 3D Protein viewer to highlight domains

CONCLUSION

- A tool for visualizing genetic variants
- Supports visualization of different levels of information
- Cross component talks
- User defined and user controlled

Google, for running the Google Summer of Code 2014.

QUESTIONS?