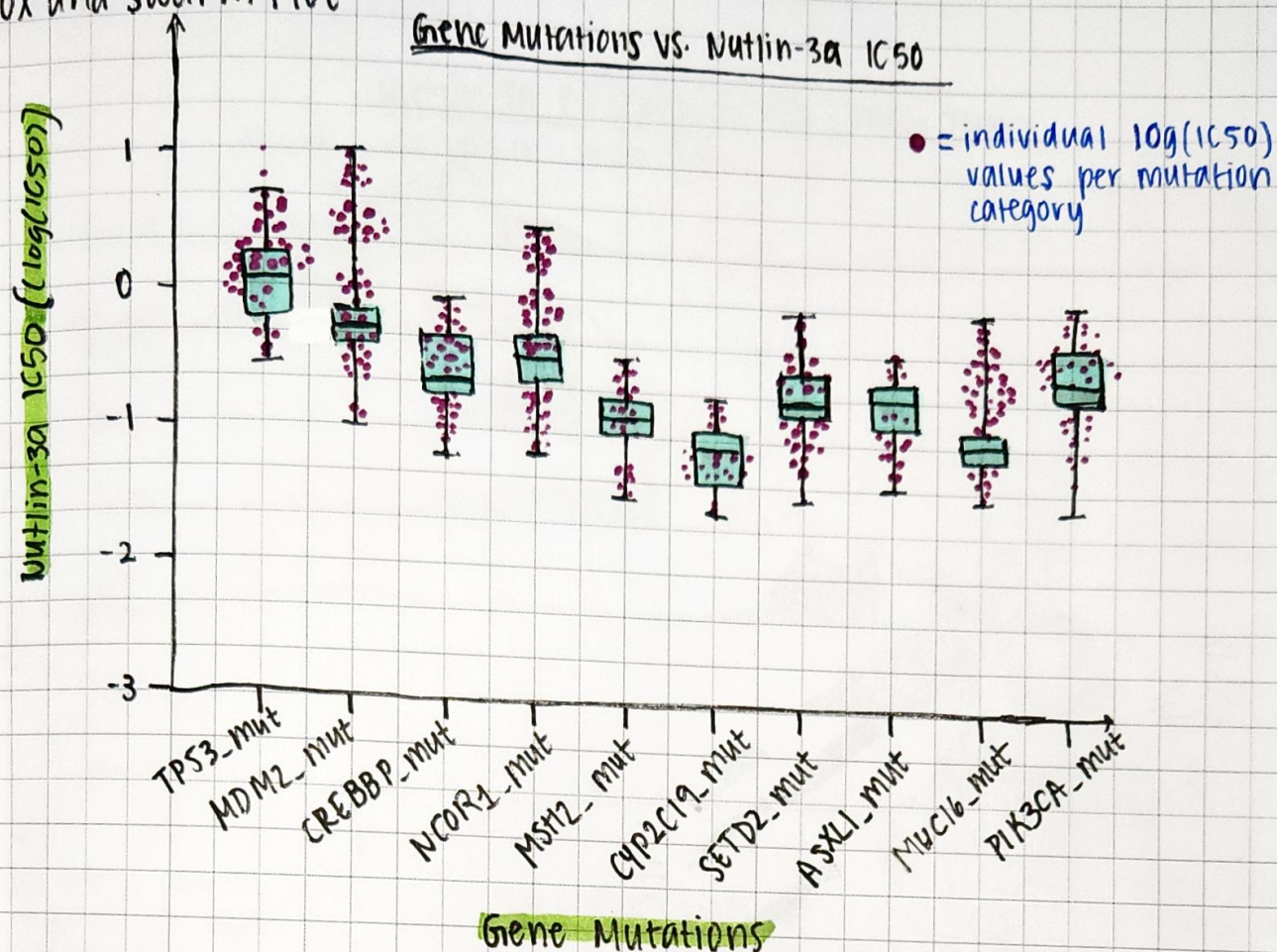
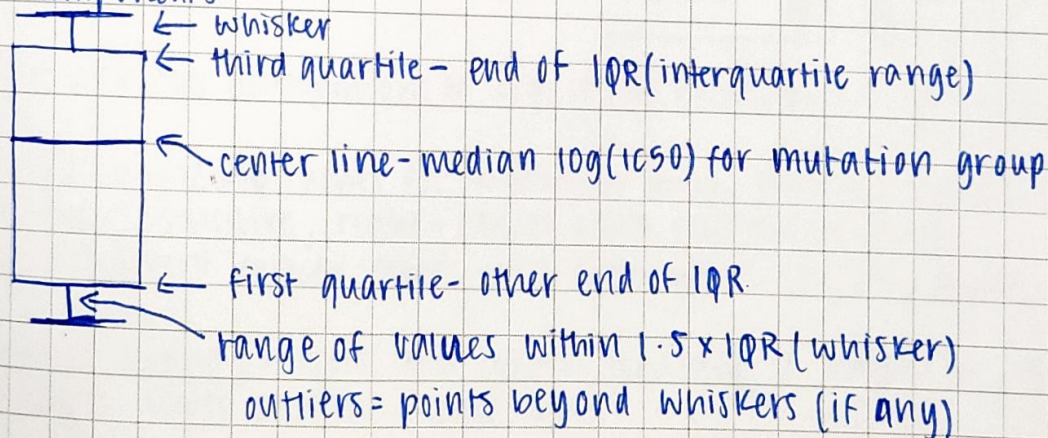


# ① Box and Swarm Plot

## Gene Mutations vs. Nutlin-3a IC50



### Box Plot components:



Swarm plots overlay individual data points; gives insight into variability & outliers. helps identify which gene mutations are associated with increased drug sensitivity/resistance as  $\text{IC}_{50}$  value directly correlates to this.

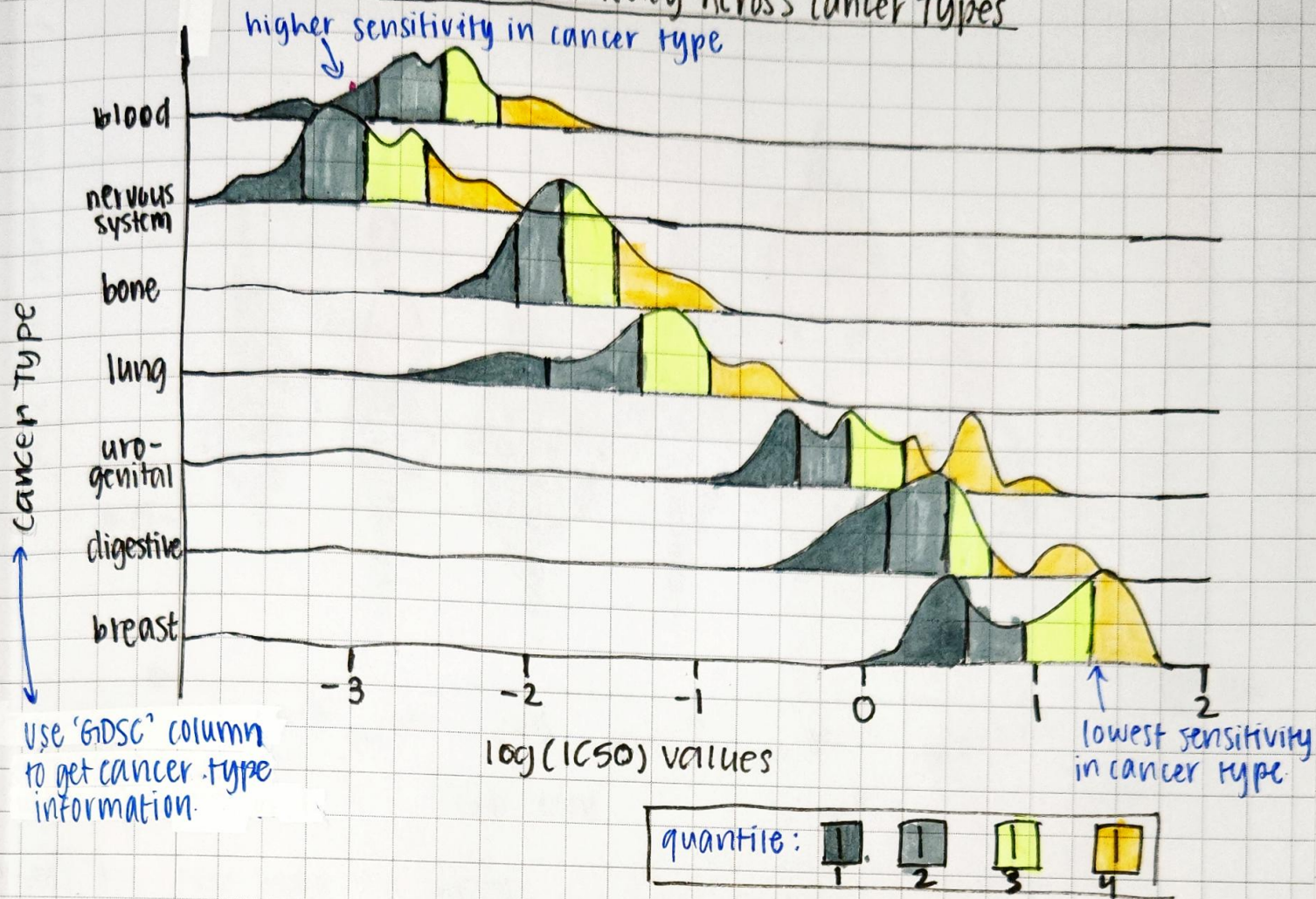
↳ low  $\text{IC}_{50}$  = less of the drug required to have an effect = higher sensitivity

data points can't overlap in swarm plot.



## ② Stacked Ridge Plot (Jogplot)

### Nutlin-3a Efficacy Across Cancer Types



Ridge plot displays distributions of  $\log(\text{IC}_{50})$  values across multiple cancer types, in this case.

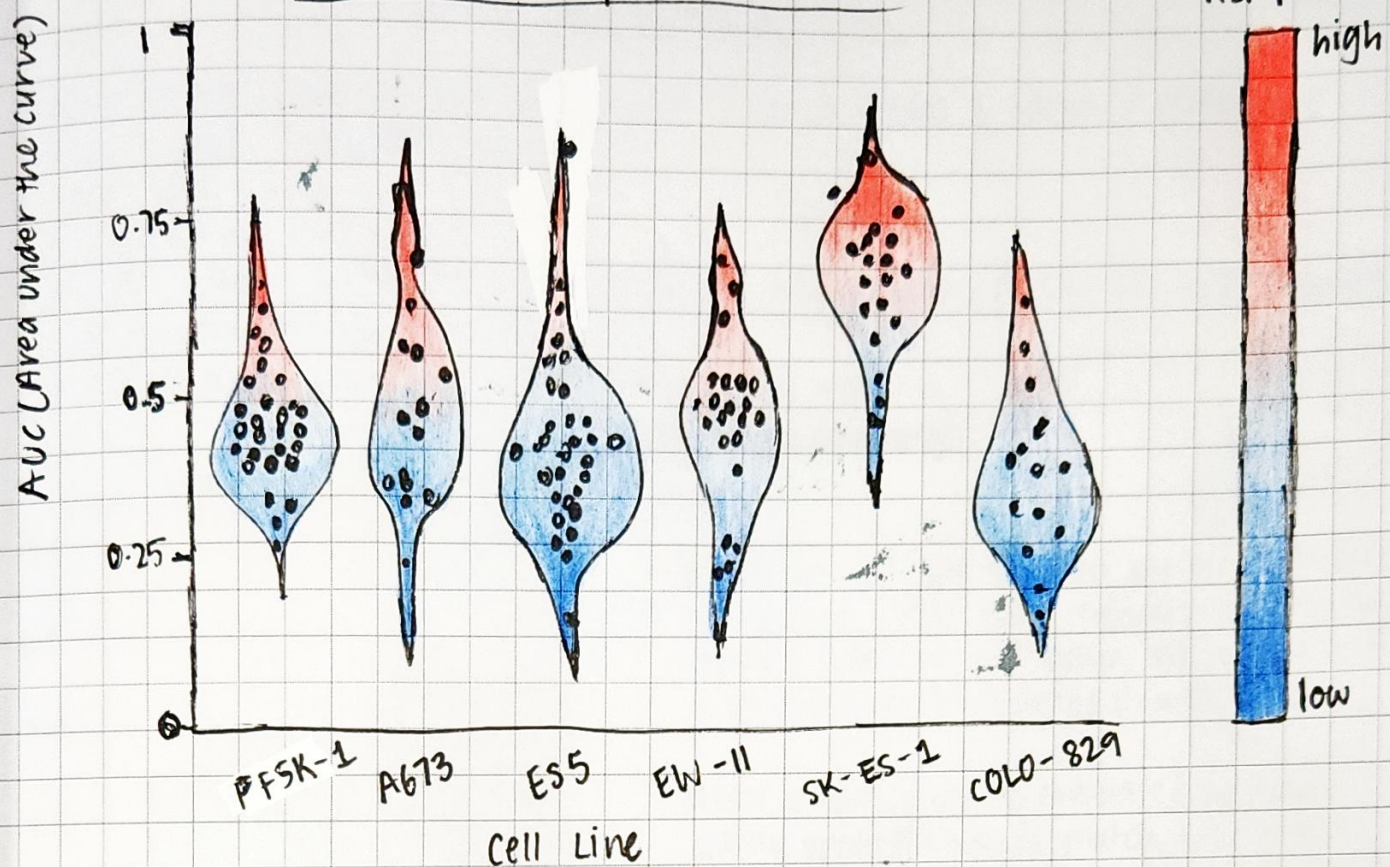
- x-axis:  $\log(\text{IC}_{50})$  values for Nutlin-3a drug, showing drug's efficacy (left = cancer more sensitive; right = cancer more resistant to drug)
- y-axis: different cancer types. each type has a smooth density curve.

can compare shape, spread, and central tendency of Nutlin-3a efficacy and see how it varies across different cancer types.



### ③ Violin Plot with strips on top:

Nutlin-3a AUC Across Cell Lines Colored by MSI p-value



Y-axis: AUC (Area under the curve)

- measures overall drug efficacy of the Nutlin-3a drug per cell line. Lower AUC = greater sensitivity (more effective). Higher AUC = greater resistance of cancer to drug (less effective). Ranges from 0 to 1.

X-axis: Cell Lines - each violin represents a unique cancer cell line. they are categorical. Violin ~~the~~ width reflects density of AUC values ~~at~~ multiple measures per cell line.

Color fill = MSI\_pval (MSI p-value) represents the statistical significance of microsatellite instability (genetic instability caused by DNA mismatch repair).

- blue = low MSI p-value → MSI-high → more genomic instability (pval < 0.05)
  - red = high MSI p-value → MSI-stable → less genomic instability (pval > 0.05)
- visually tracks relationship between MSI & AUC.

Individual points for better specificity (with the black color). data points overlap allowed.

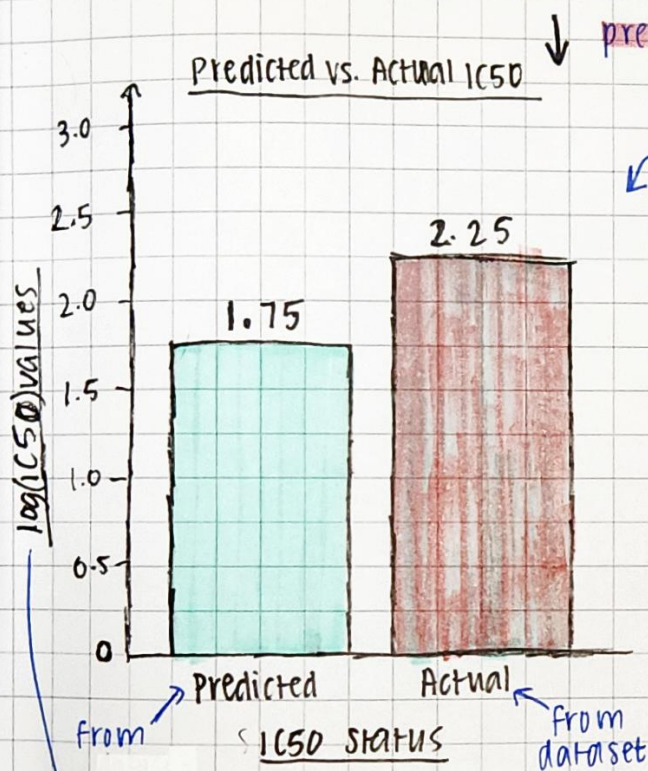


## ④ Machine Learning Model

### Predicting Nutlin-3a IC50 from Biological Features

Input values / features (that will be used to train model & identify patterns):

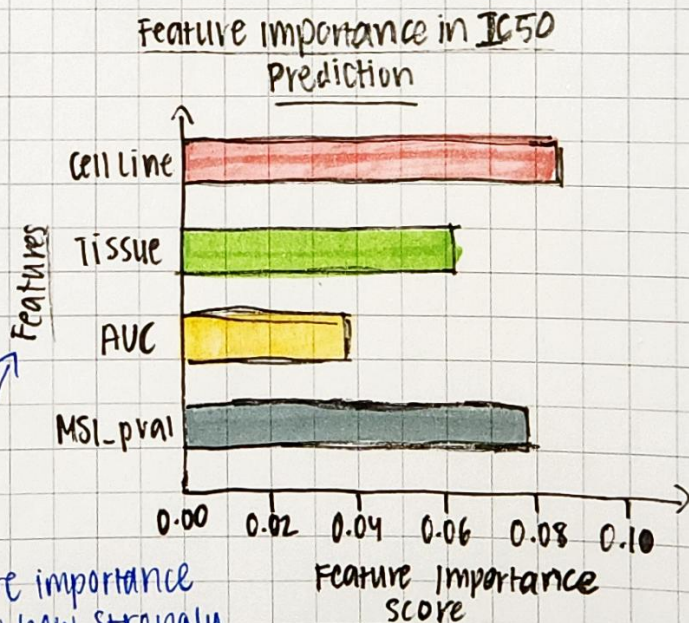
- MSI - pval (MSI p-value)
- Cell Line Name
- GDSC (tissue classification)
- ic50 - effect - size (measures difference in drug response (IC50) between mutated & nonmutated samples)
- AUC or area under the curve: overall drug response.



↓ predicts IC50 efficacy

Nutlin-3a efficacy based on various features is predicted. Other drug features can also be inputted (that are not Nutlin-3a) so that IC50 is predicted for other drugs as well.

Model accuracy can be evaluated by how closely predicted values match true IC50 values. Larger gaps indicate error.



plotting log values (for better interpretability) but the original IC50 is measured in  $\mu\text{M}$ .

Feature importance scores show how strongly each feature influences the predicted IC50 values. Higher scores = greater predictive power different for each kind of model.



## ⑤ Interactive hypothetical prediction simulator

### Simulate NUTlin-3a Response for Hypothetical Patients

#### Patient Features

adjusts  
microsatellite  
instability p-  
value. Shows  
how unstable  
the genome  
is.

◦ MSI (microsatellite instability)  
p-value



◦ Tissue Type

can select  
tissue type  
(liver, lung,  
breast, etc.)

Lung V

◦ Molecular Features

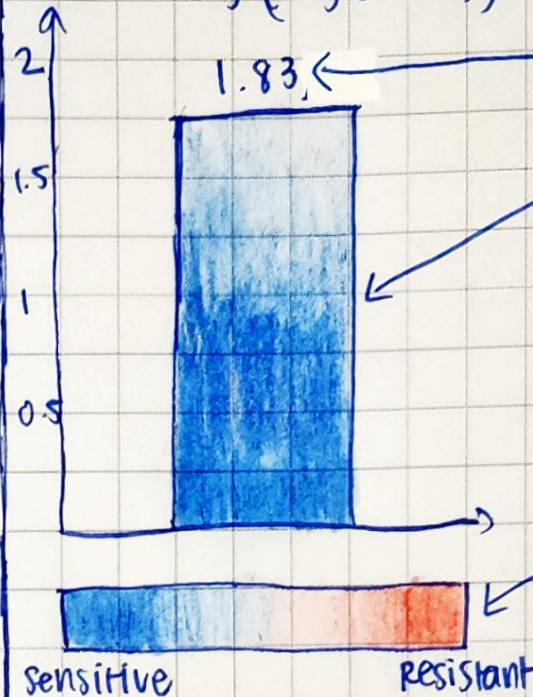
☒ TP53 Expression-high

☒ MDM2 upregulation

☐ CDKN2A deletion

switches for  
key biomarkers  
on/off status

#### Predicted NUTlin-3a Efficacy ( $\log(IC_{50})$ )



lower =  
more  
effective  
dynamic  
updates to  
show  
NUTlin-3a  
response  
based on  
selected  
features  
sensitivity  
of cancer  
cells to  
NUTlin-3a