

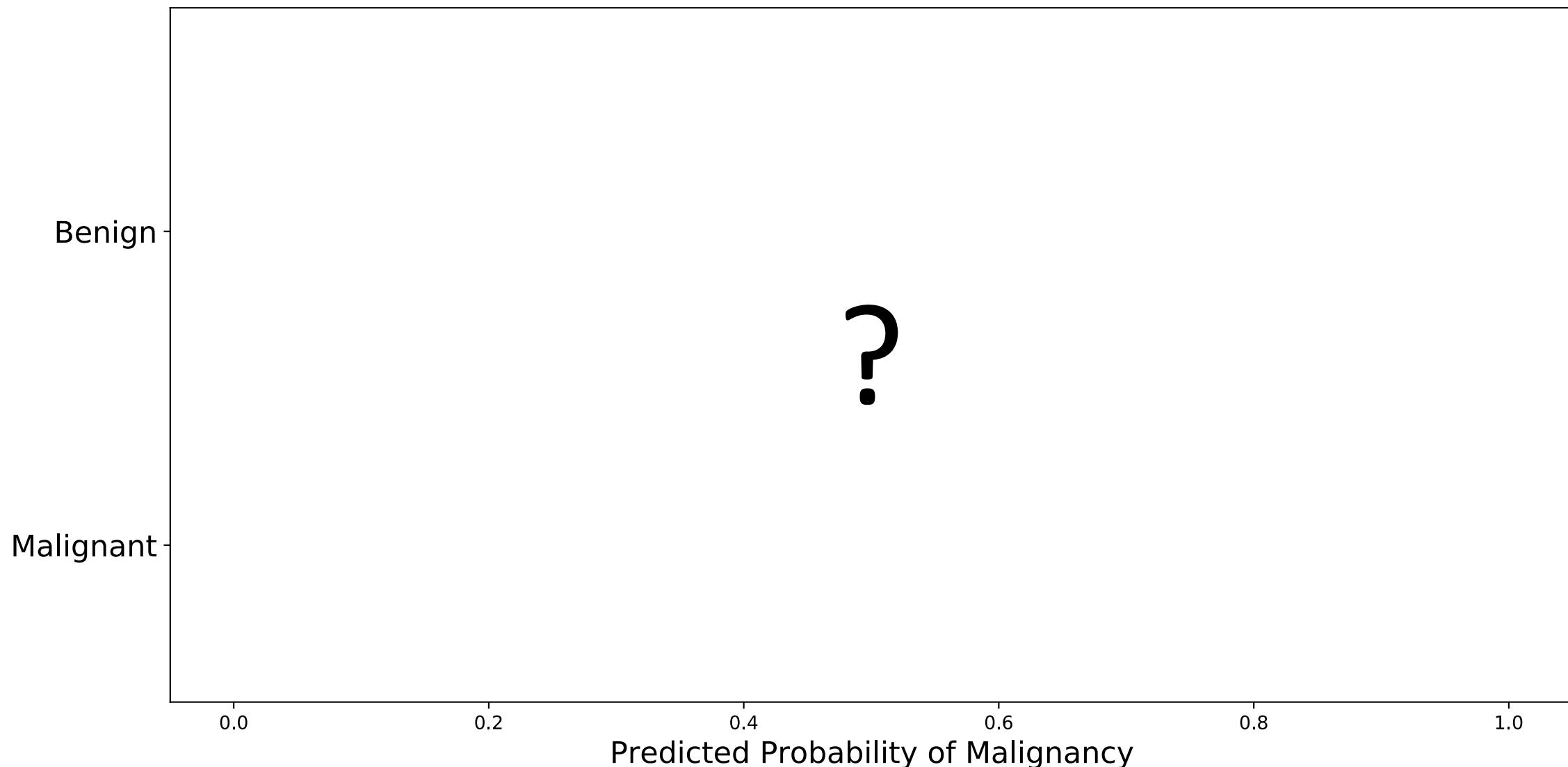
# Performance Measures

Matthew Engelhard

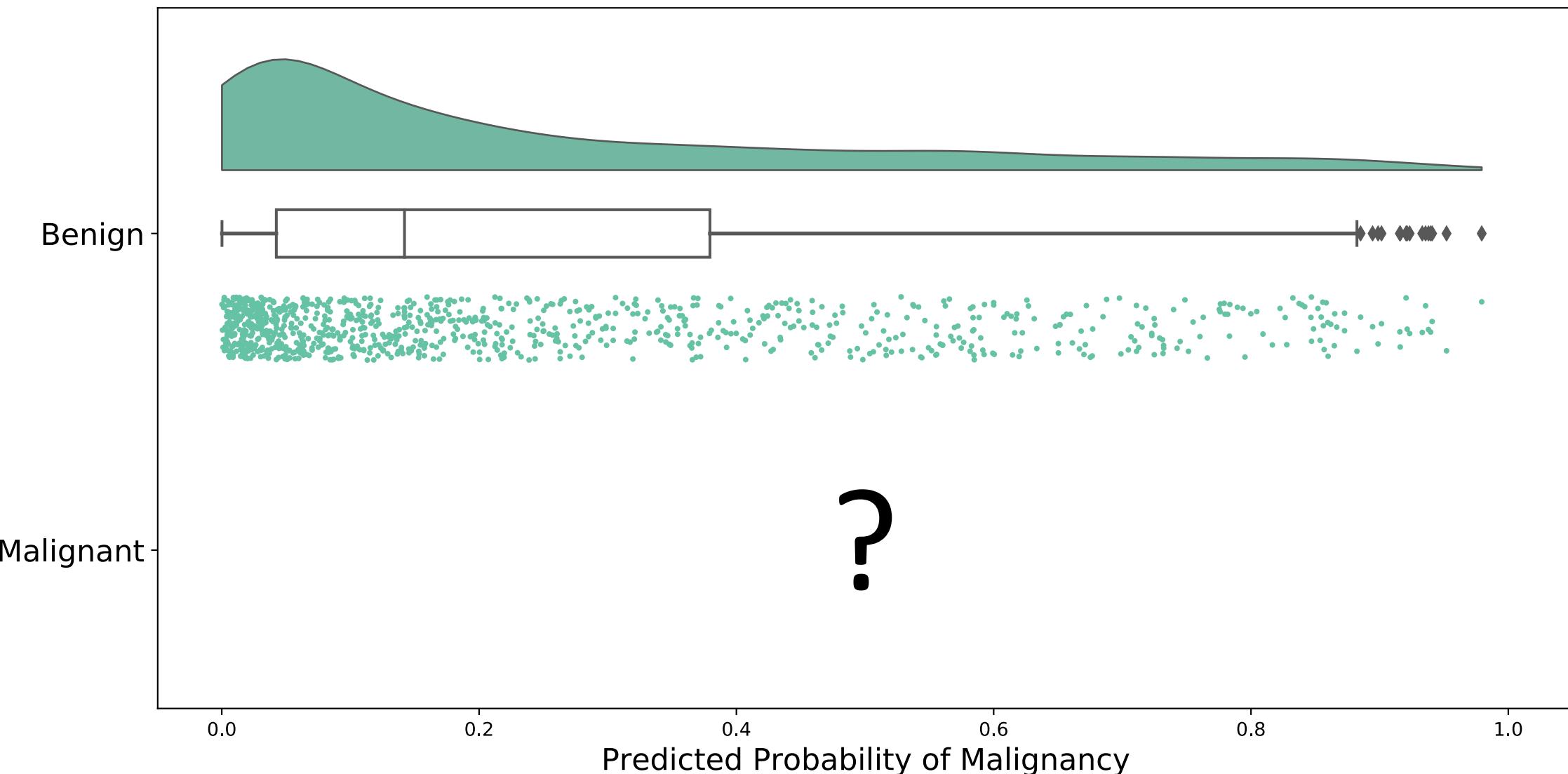
# Goals

- Understand and calculate common performance measures for binary classification
- Contextualize performance against that of a *no information* classifier
- Recognize that *good* performance depends on existing alternatives
- Match clinical scenarios to performance measures important in that scenario

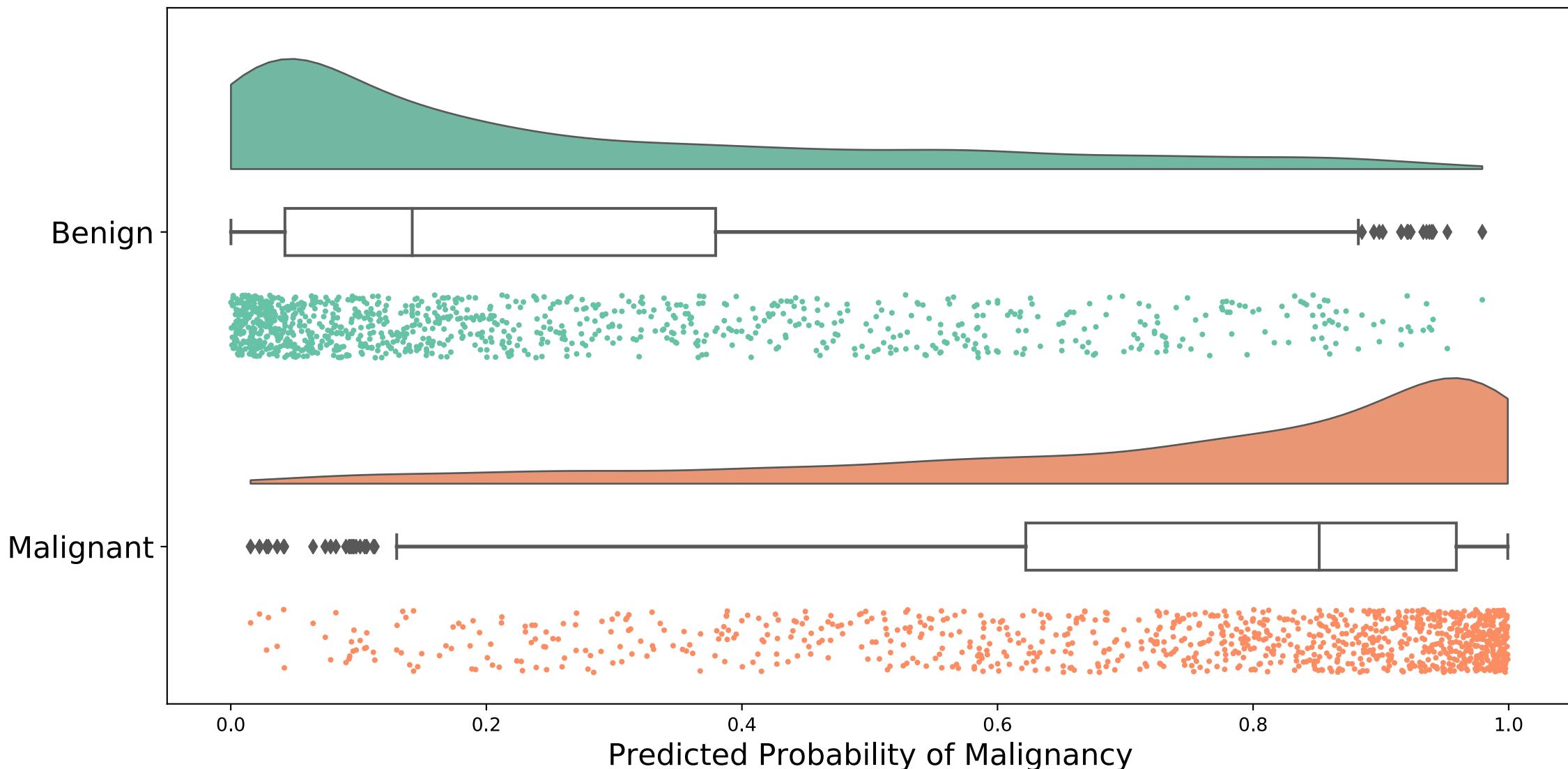
Back to cancer prediction. Suppose our features are highly informative. What might our model's predictions look like?

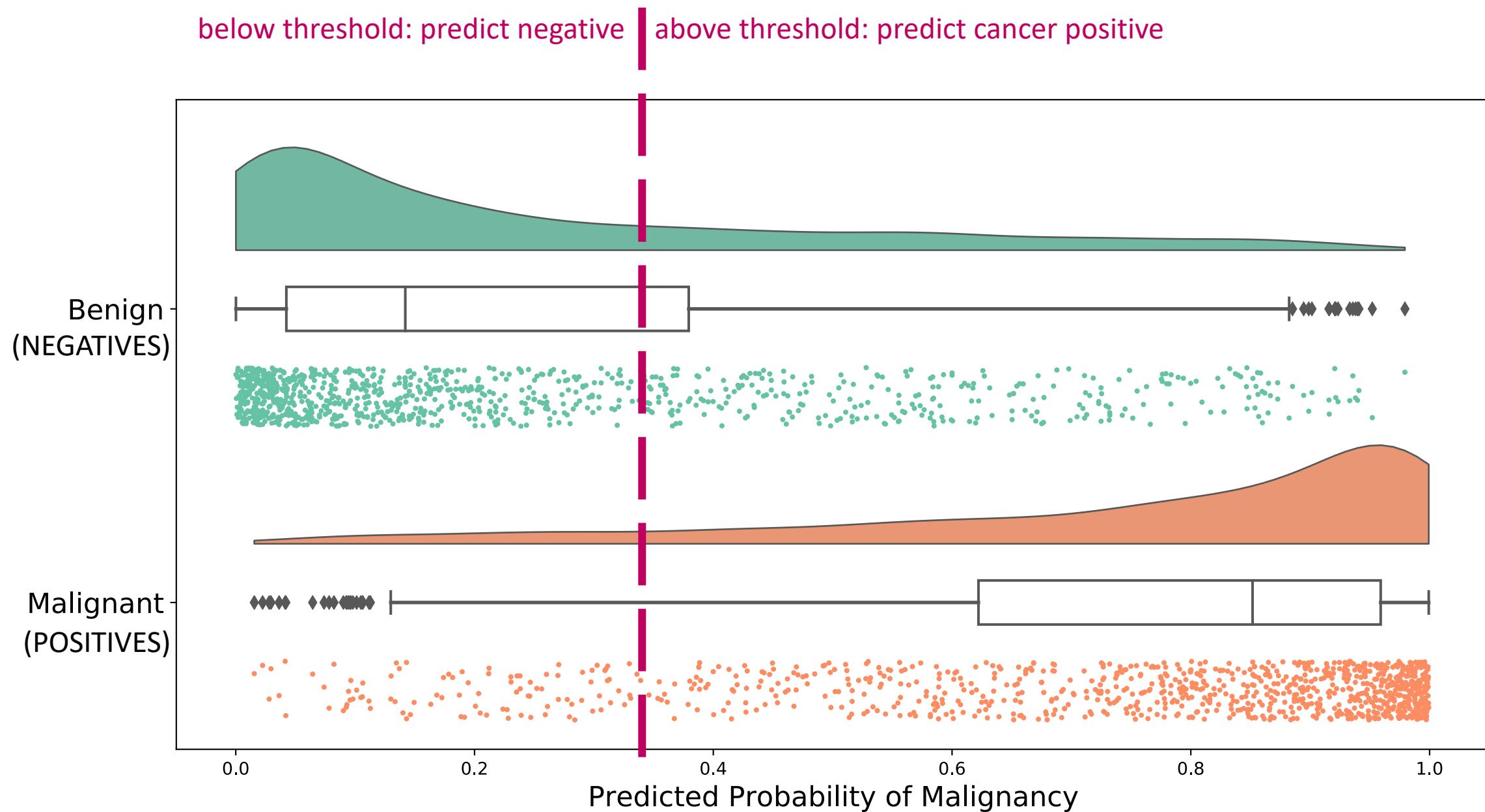


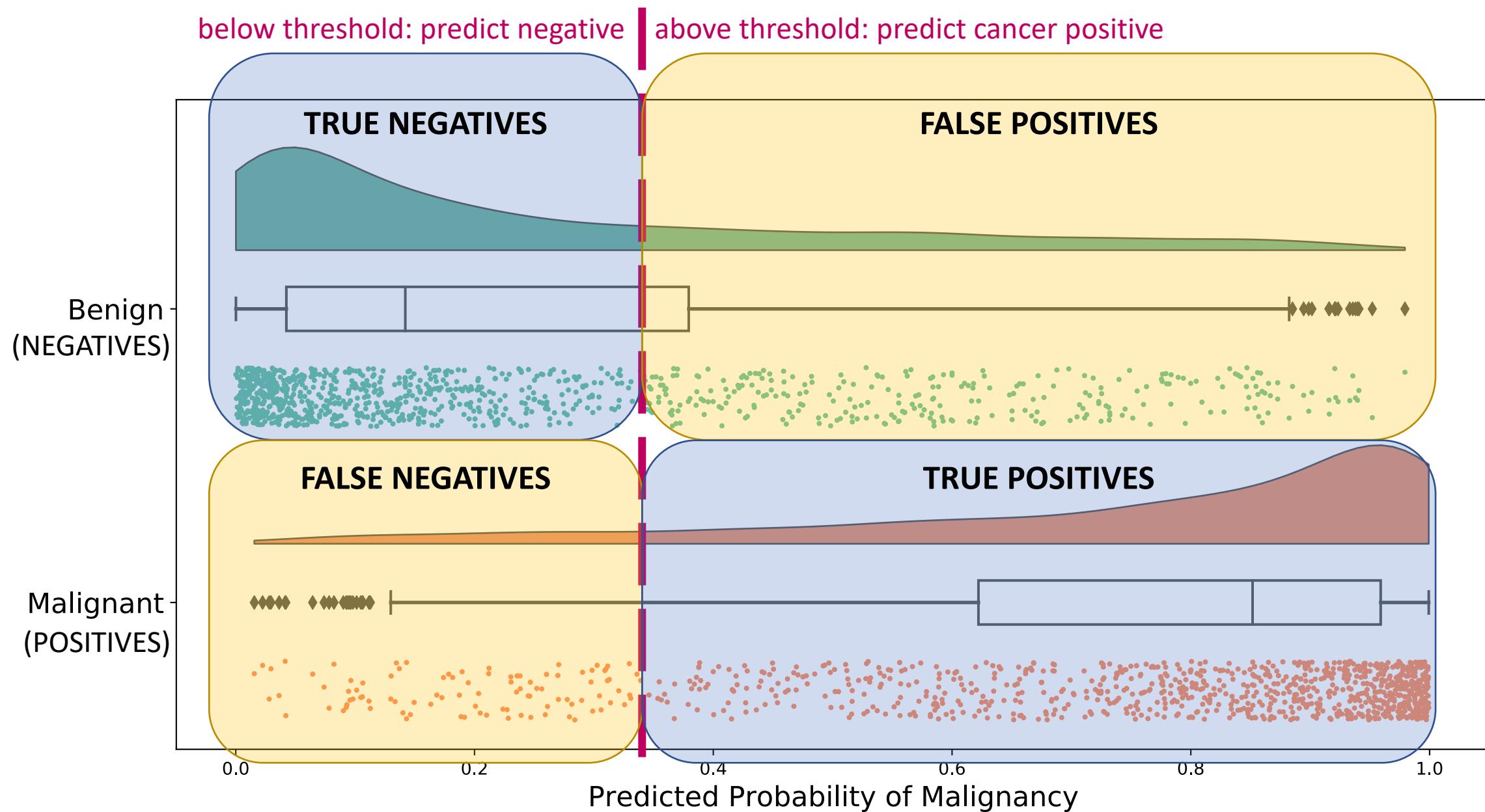
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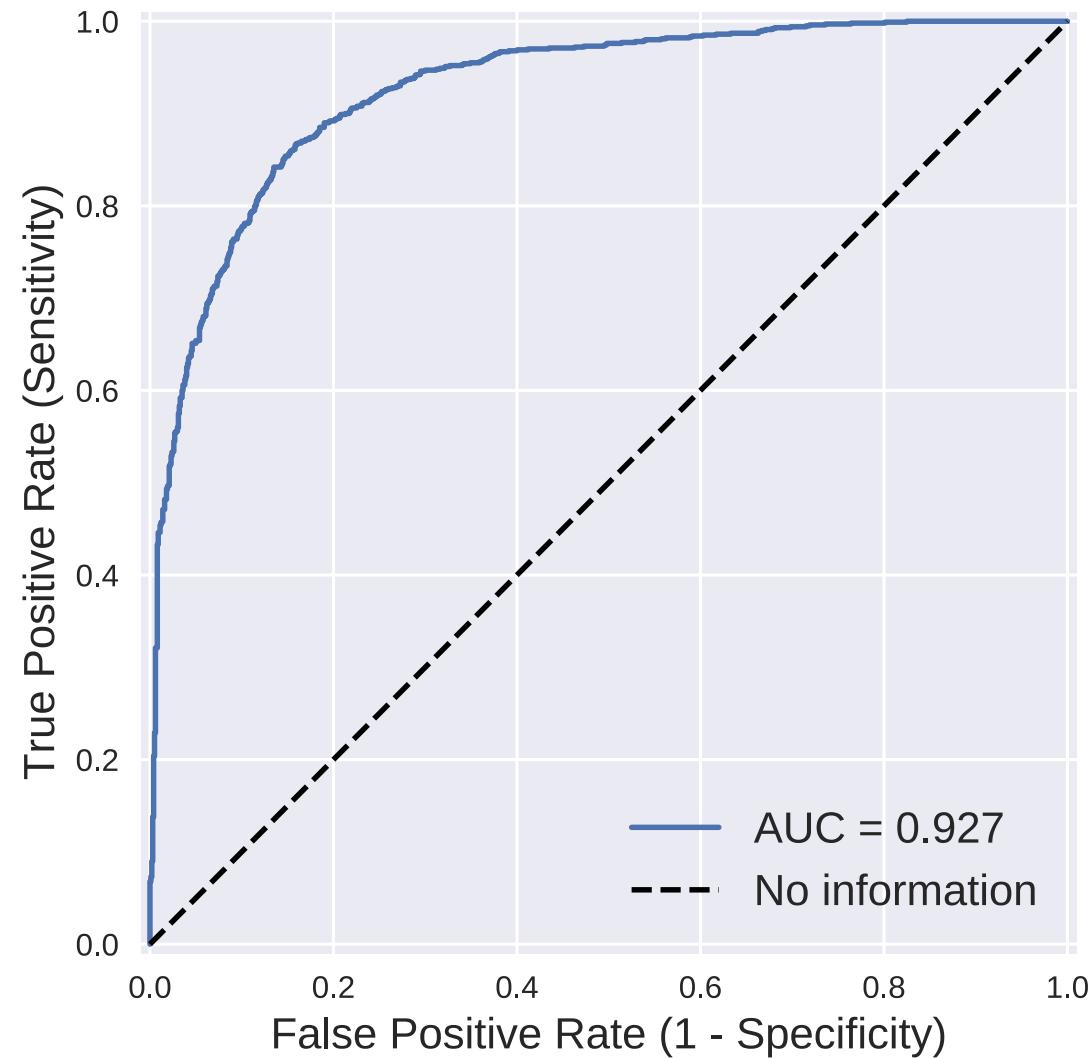






# Receiver Operating Characteristic Curve

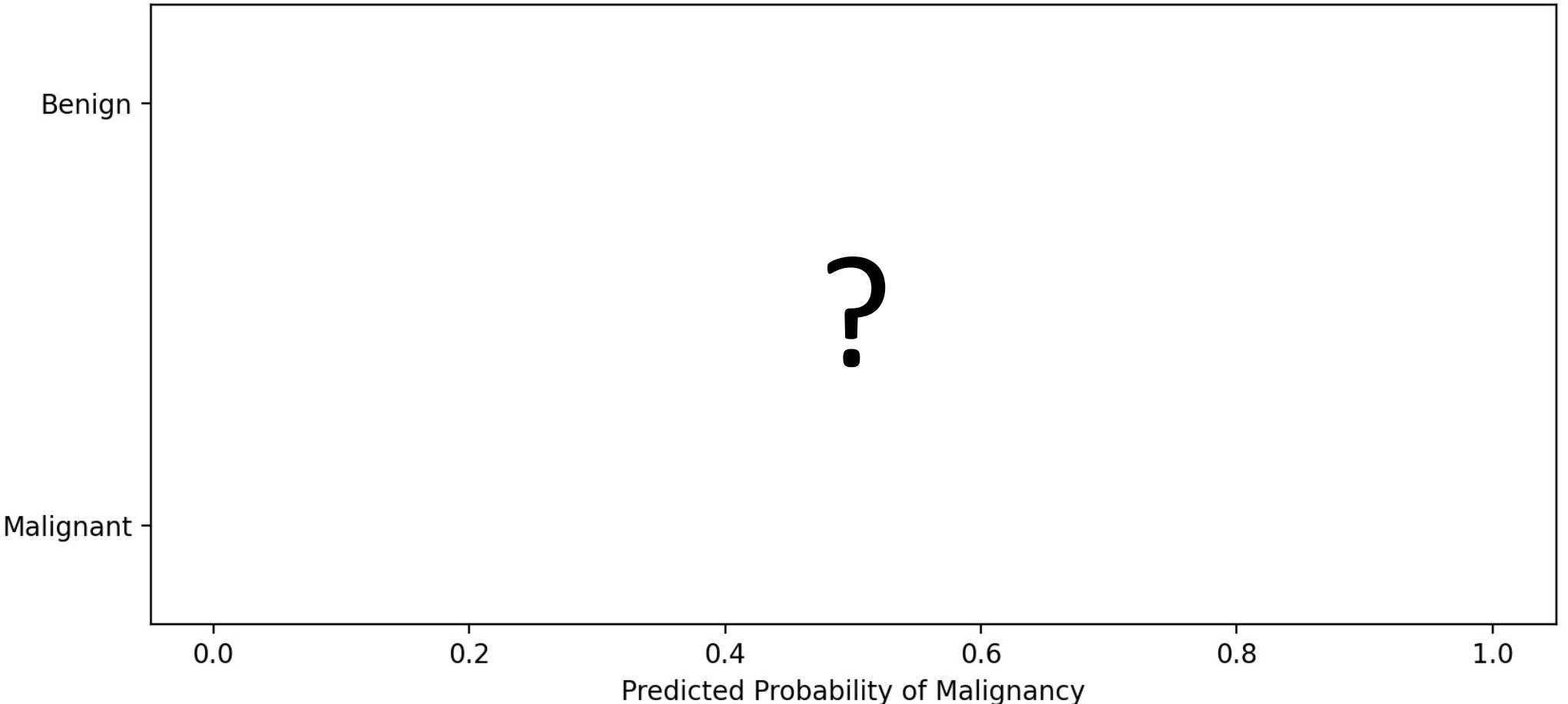
- Illustrates the tradeoff between the true positive rate (i.e., sensitivity) and the false positive rate (i.e.,  $1 - \text{specificity}$ ) as we vary the threshold.
- The area under this curve (AUC) provides a single summarizing this tradeoff.
- Note that to get the sensitivity versus specificity curve, we simply rotate the ROC curve clockwise by 90 degrees. The areas under the two curves are the same.



So, what's a *good* AUC value?  
(i.e., *good* performance)?

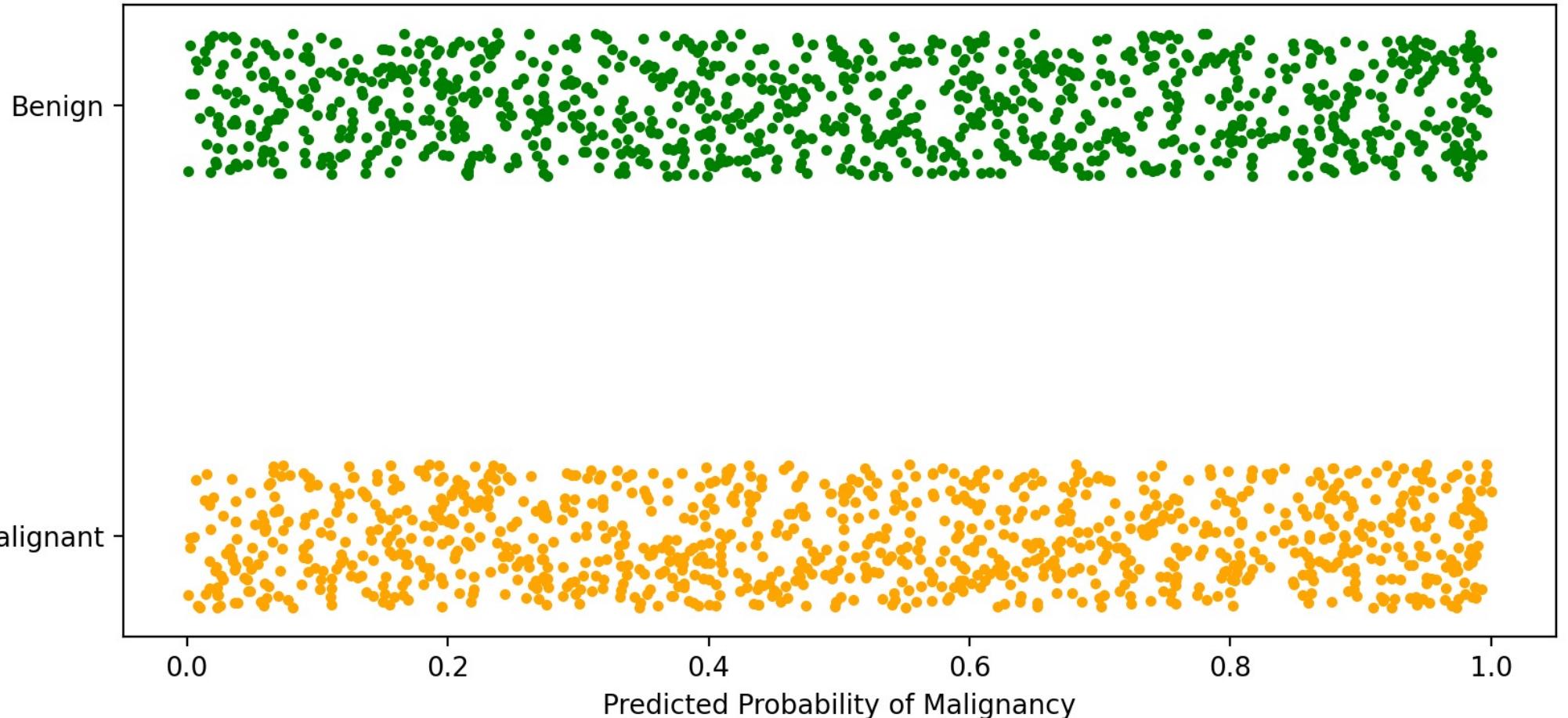
We'll start to answer this question by taking a look at *bad* performance.

Suppose our features contain *no information* about the label.  
What might our model's predictions look like?



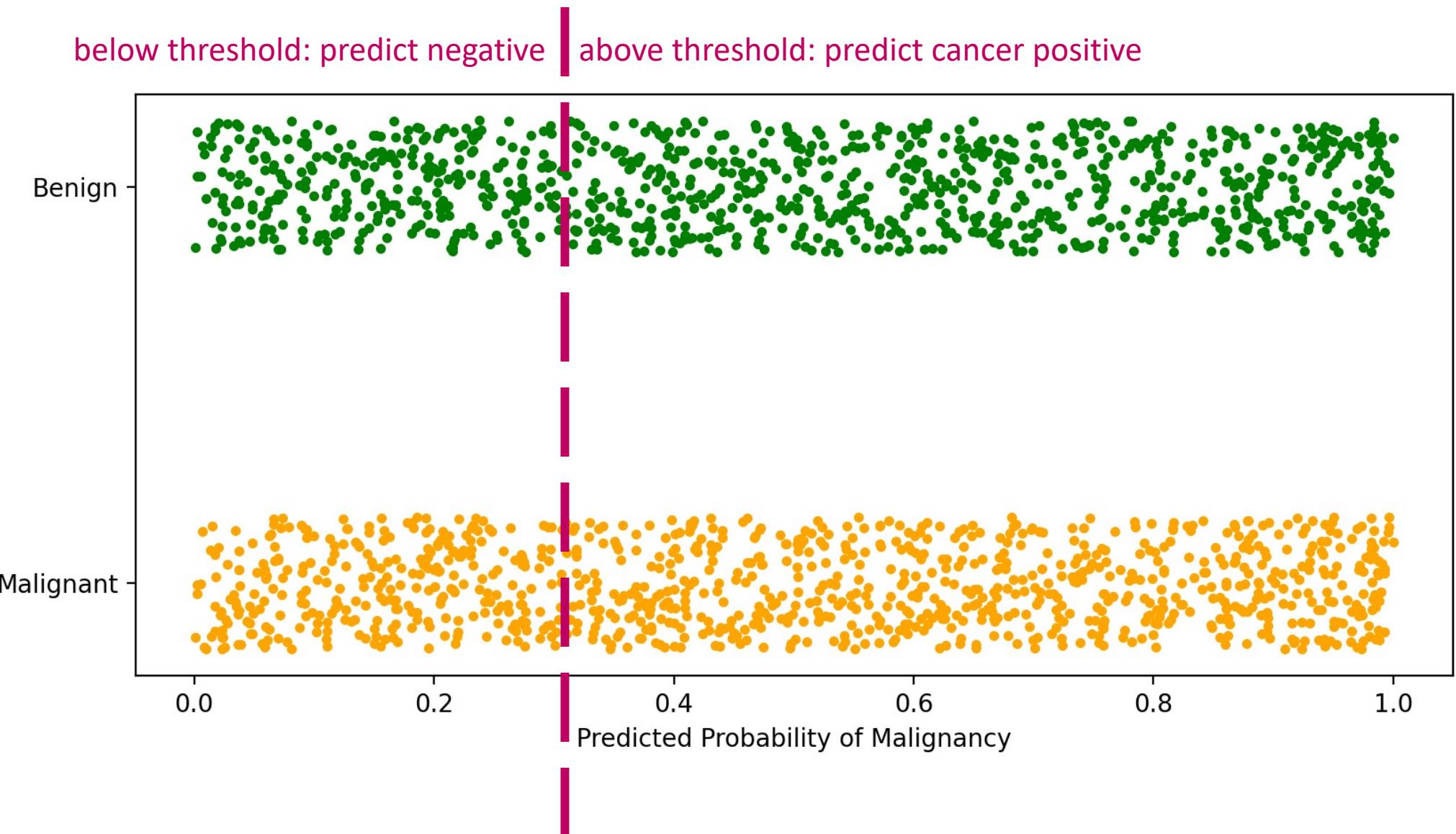
Suppose our features contain no information about the label.  
What might our model's predictions look like?

- Similar distributions between positive and negative cases.
- The predicted value tells you nothing about which one it's more likely to be.



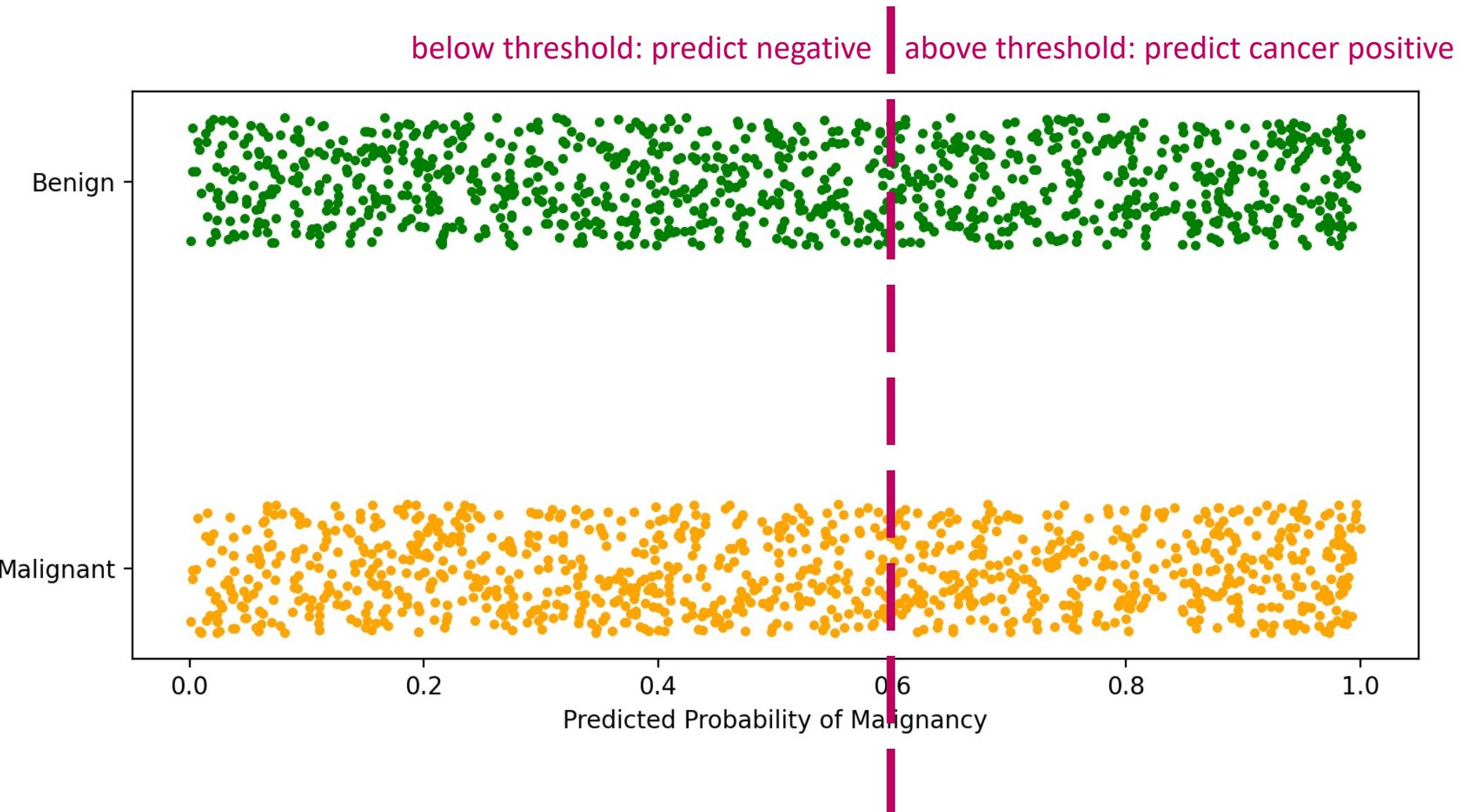
# We'll try placing a threshold just like before

What is the:  
(a) Sensitivity?  
(b) Specificity?  
(c) Positive  
predictive  
value?



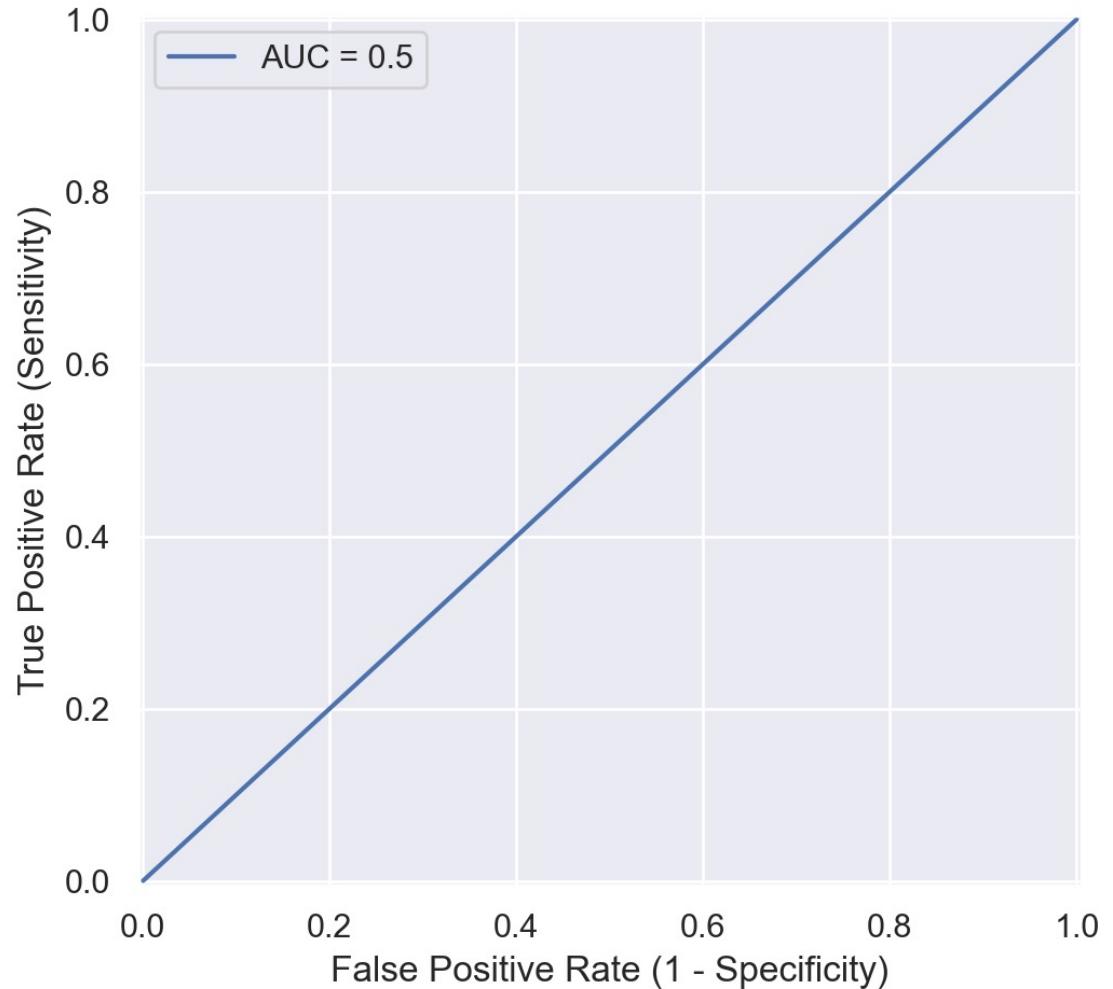
# We'll try placing a threshold just like before

What is the:  
(a) Sensitivity?  
(b) Specificity?  
(c) Positive  
predictive  
value?



# *Our no information predictive model:*

- Place threshold at .3
  - Sensitivity = .7
  - False positive rate = .7
  - Specificity = 1-.7
- Place threshold at  $p$ 
  - Sensitivity =  $1-p$
  - False positive rate =  $1-p$
  - Specificity =  $p$



# Let's think about it a different way.

- Suppose we have no predictors – again, no information – so we decide we'll just flip a coin instead of building a model.
  - If the coin comes up *heads*, we'll predict *positive*.
  - If the coin comes up *tails*, we'll predict *negative*.



Fair Coin:  $P(\text{heads}) = .5$

- Sensitivity = ?
- False positive rate = ?
- Specificity = ?

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Fair Coin:  $P(\text{heads}) = .5$

- Sensitivity = .5
- False positive rate = .5
- Specificity = .5



Biased Coin:  $P(\text{heads}) = p$

- Sensitivity = ?
- False positive rate = ?
- Specificity = ?

# Let's think about it a different way.

- Suppose we have no predictors – again, no information – so we decide we'll just flip a coin instead of building a model.
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Fair Coin:  $P(\text{heads}) = .5$

- Sensitivity = .5
- False positive rate = .5
- Specificity = .5

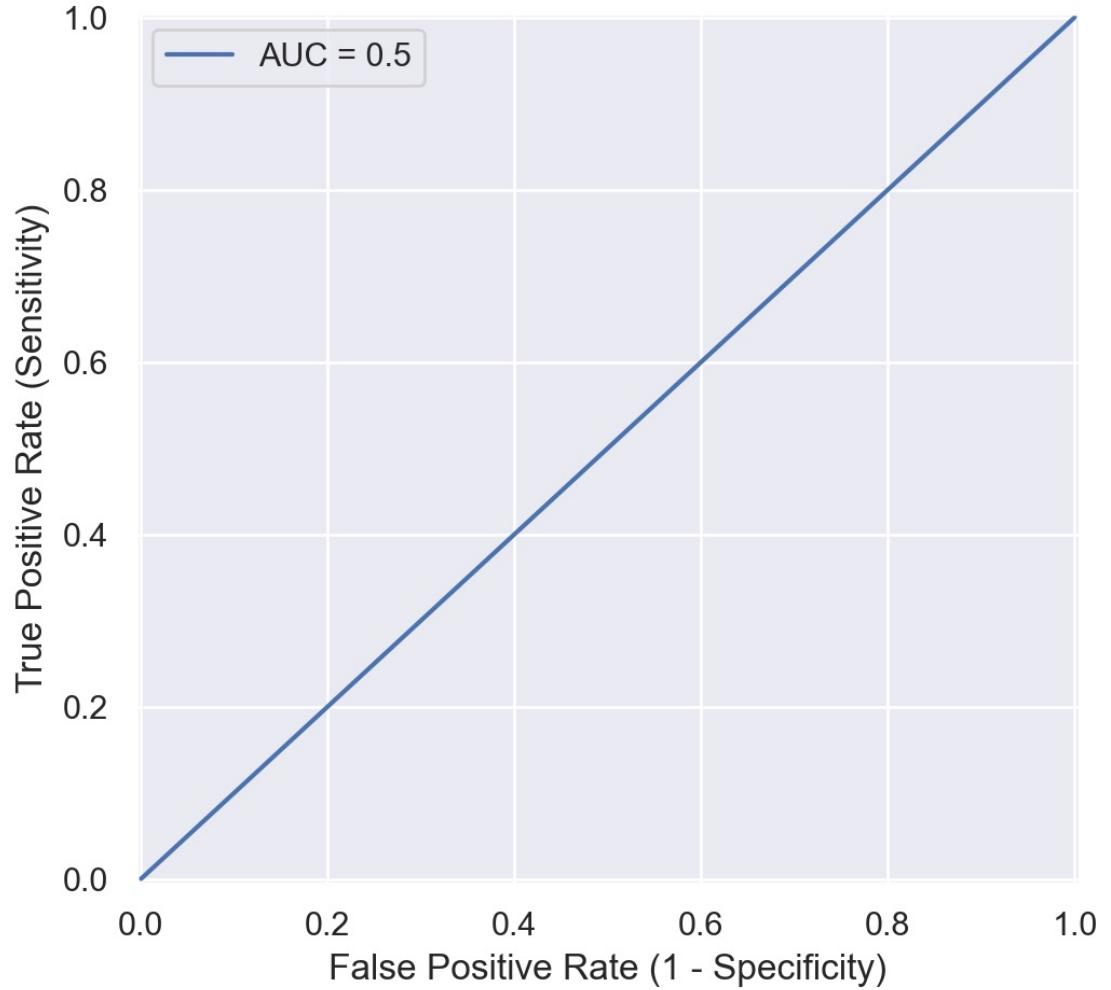


Biased Coin:  $P(\text{heads}) = p$

- Sensitivity =  $1-p$
- False positive rate =  $1-p$
- Specificity =  $p$

Again, we arrive at the following *no information* curve

- We may choose any  $p$  between 0 and 1 to get:
  - Sensitivity =  $p$
  - False positive rate =  $p$
  - Specificity =  $1-p$
- What's the area under this curve (AUC)? --> 0.5



So, what's a *good* AUC value?  
(i.e., *good* performance)?

It depends.

- Are predictions better than random?
- Are predictions than the previous best performing model?
- Are predictions better than expert performance?
- Does performance exceed our (informed) expectations?
- **Is the model clinically useful?**

# OK, we've quantified performance across all thresholds. But how do we use the model?

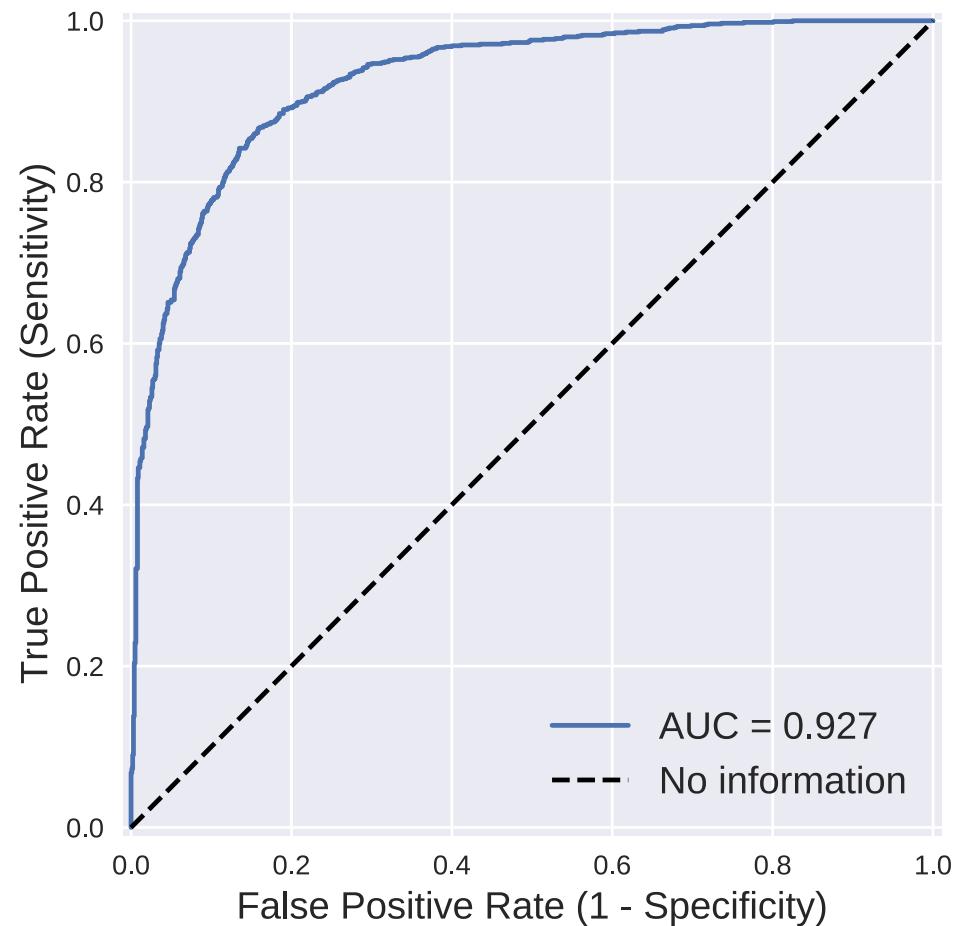
Sometimes the predicted probability really is what we care about.

- *Example*: probability of heart attack
- If so, we need to make sure our model is *calibrated*

More often, we need to pick a threshold so we can decide whether to:

- Alert a provider
- Get a biopsy
- Refer the patient
- etc

What threshold should we pick? What's the right tradeoff?



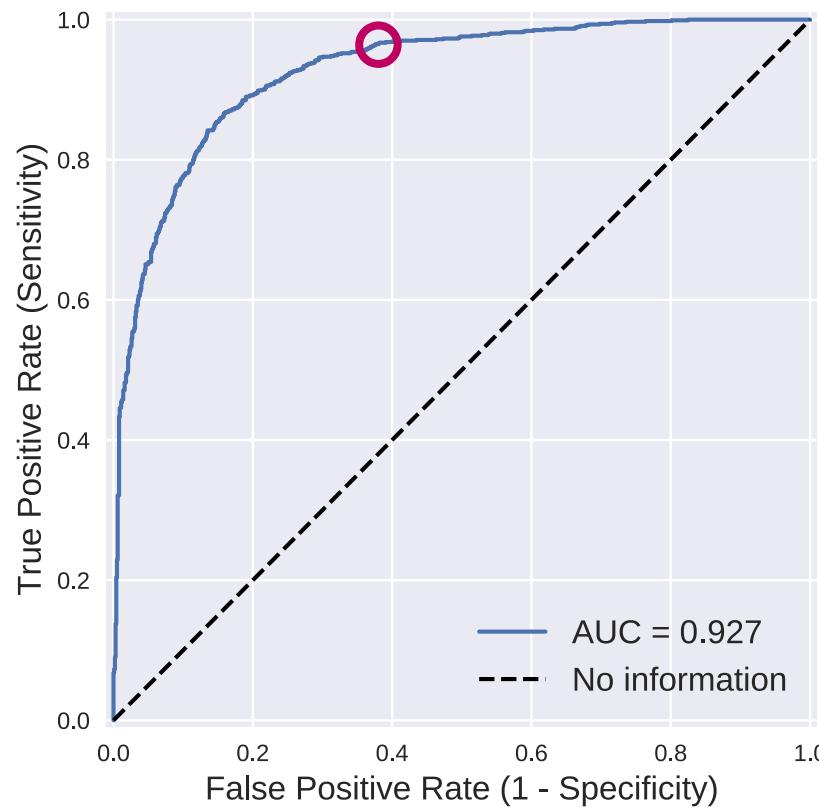
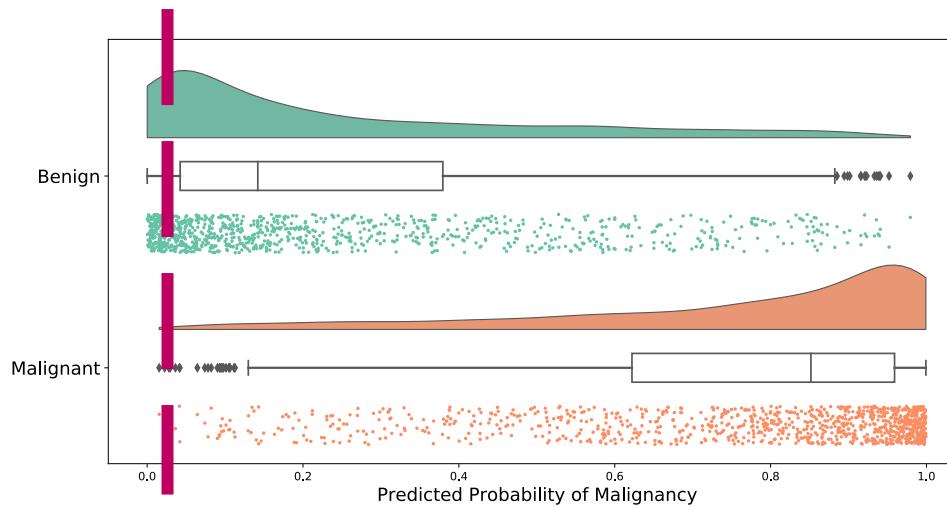
# Healthcare Scenarios

*Which performance measure is most important?*

1. A computer vision model that detects carcinoma

# Operating Point:

*high sensitivity*

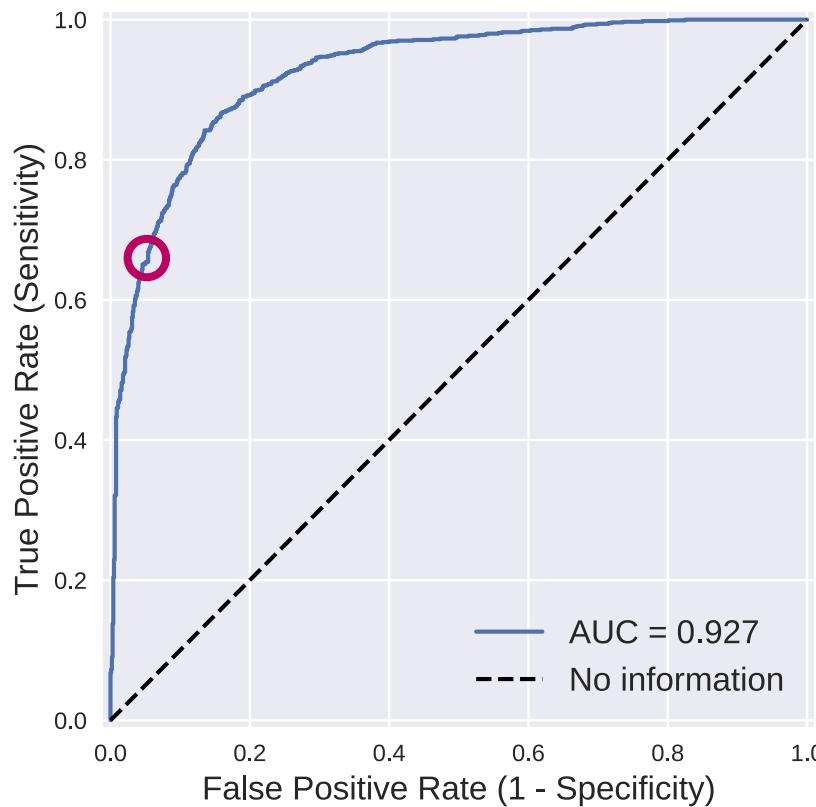
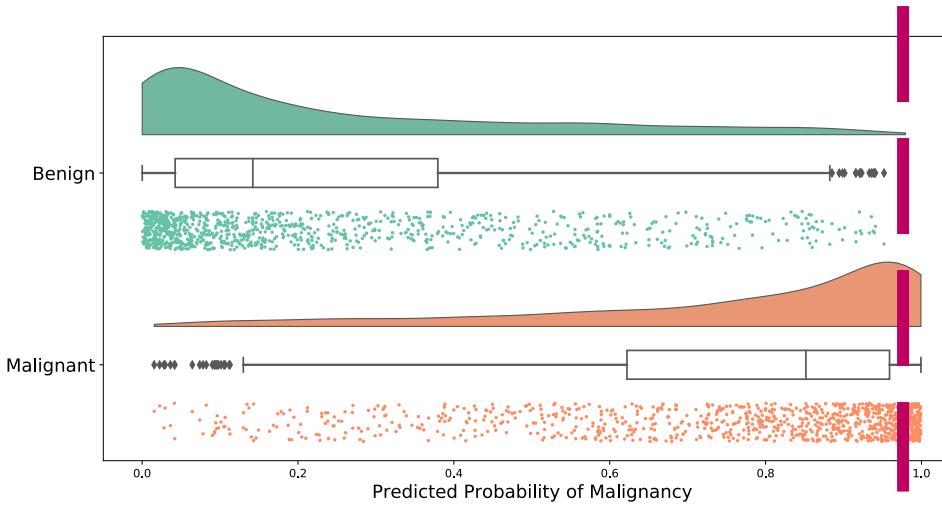


# Healthcare Scenarios

1. A computer vision model that detects carcinoma
2. An algorithm that detects atrial fibrillation in Apple Watch users

# Operating Point:

*high specificity*

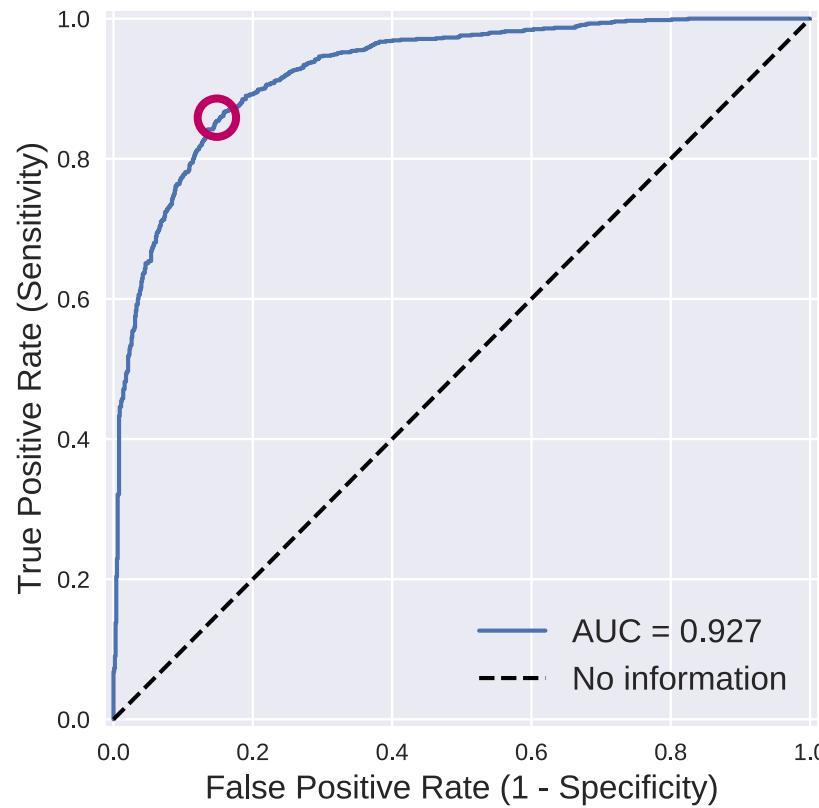
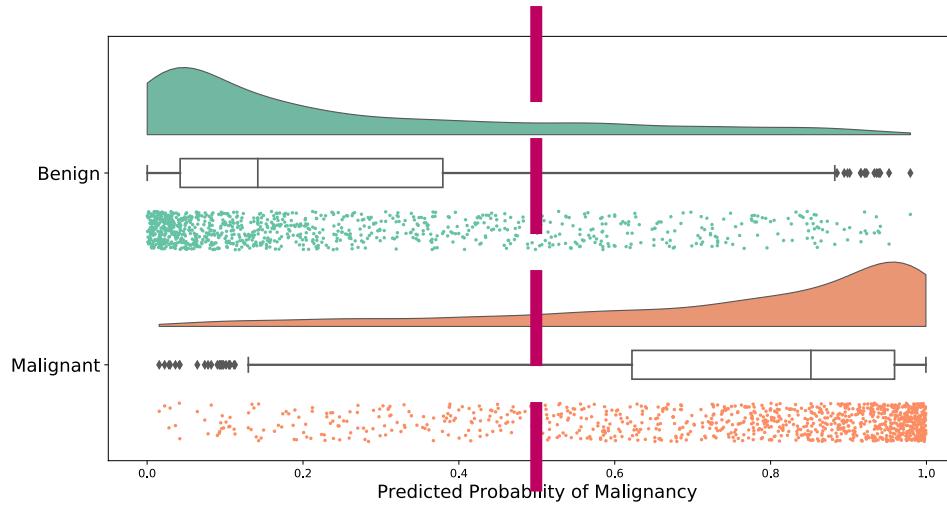


# Healthcare Scenarios

1. A computer vision model that detects carcinoma
2. An algorithm that detects atrial fibrillation in Apple Watch users
3. An EHR-based model that monitors autism risk

# Operating Point:

*balanced*



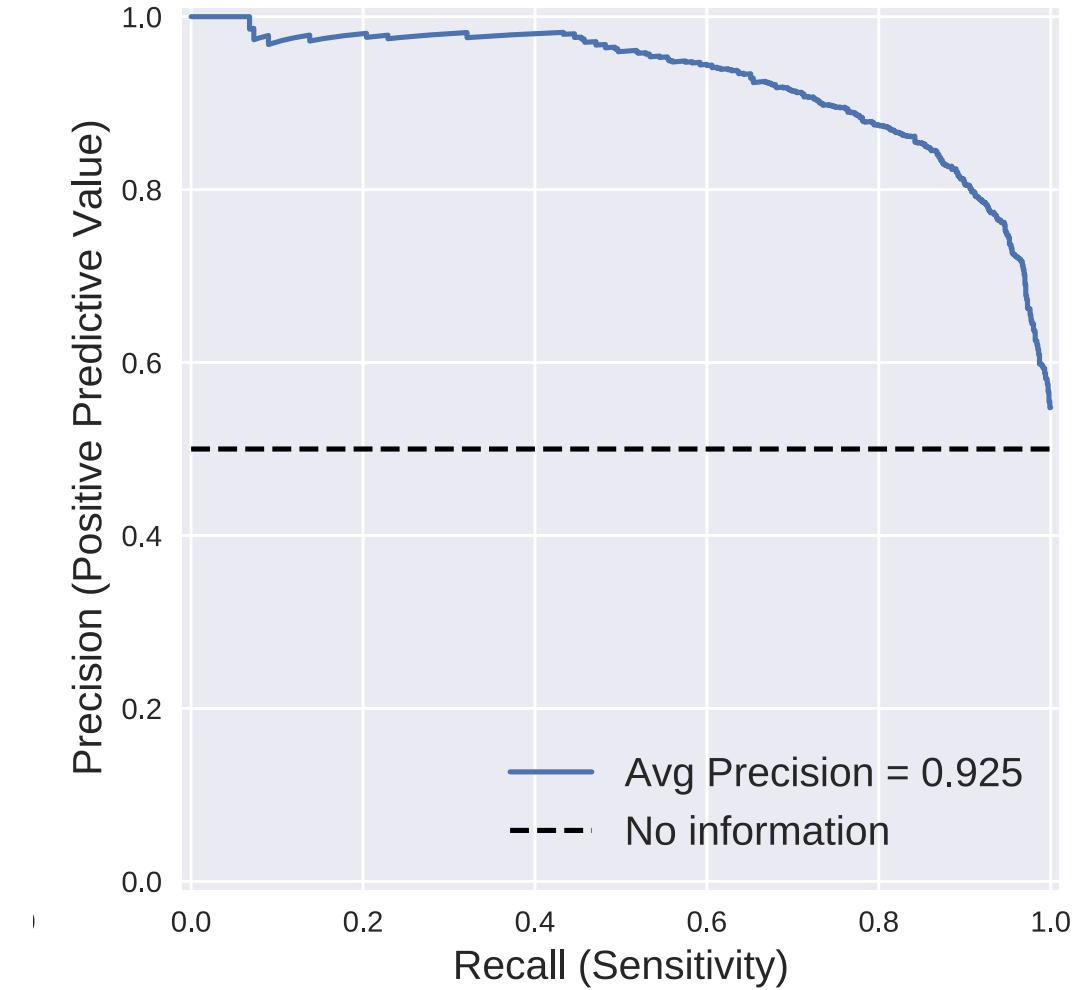
# Healthcare Scenarios

1. A computer vision model that detects carcinoma
  2. An algorithm that detects atrial fibrillation in Apple Watch users
  3. An EHR-based model that monitors autism risk
- 
- Sometimes specificity and sensitivity are difficult to interpret, particularly for rare conditions or events.
  - The most clinically relevant measure is often the positive predictive value (or negative predictive value).

# The Precision-Recall Curve

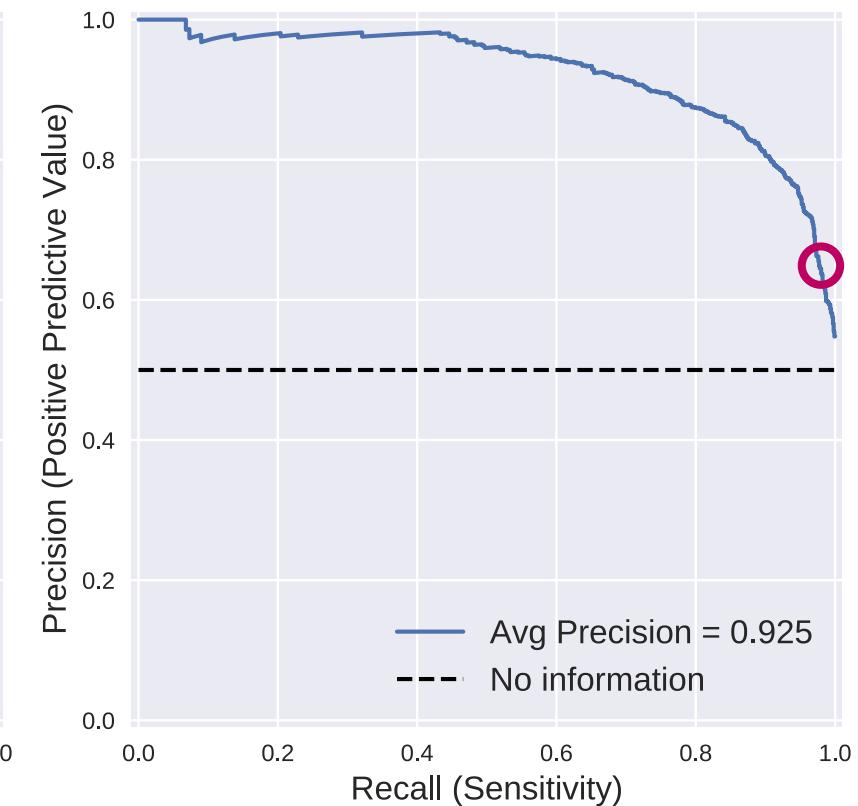
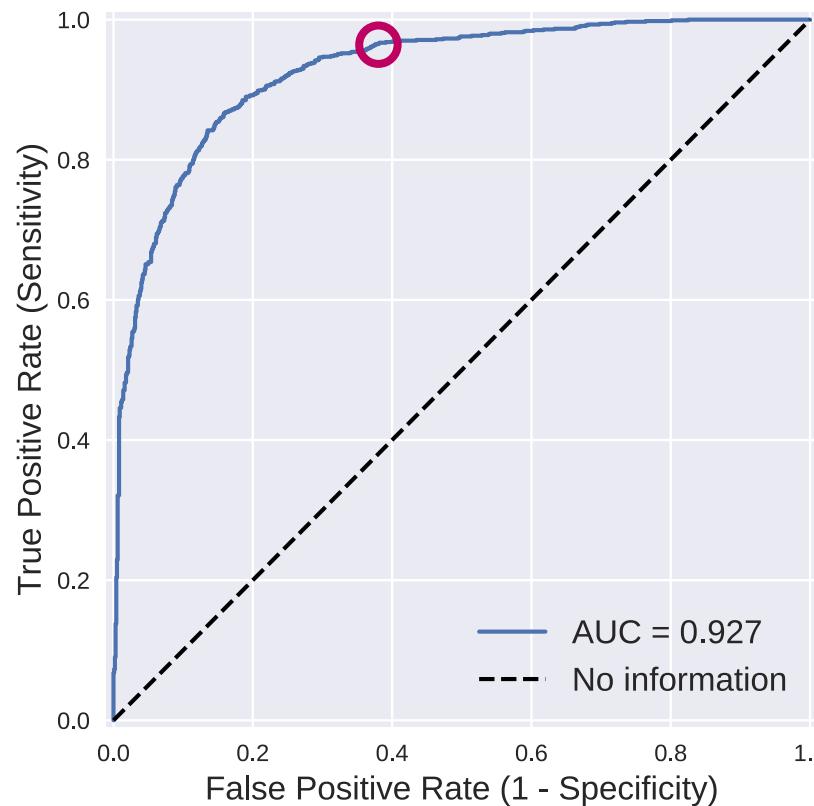
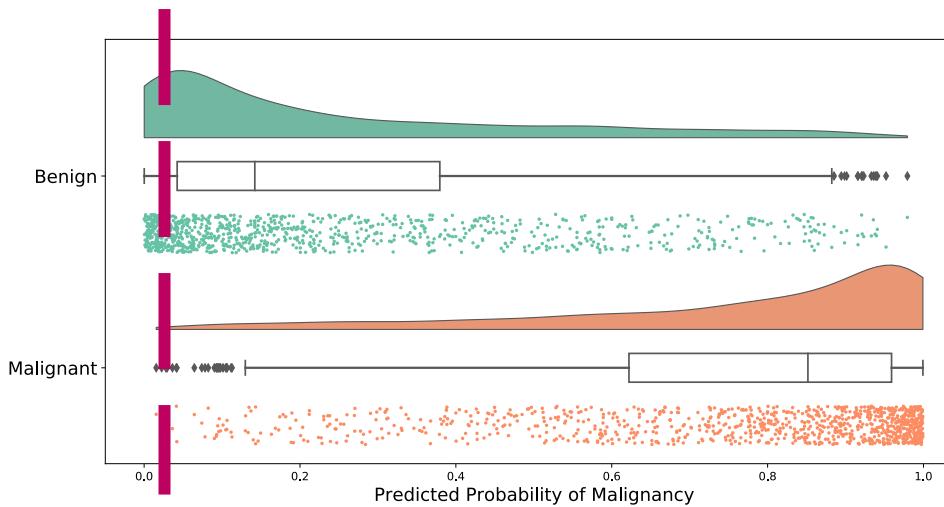
(i.e., PPV-Sensitivity Curve)

- Often has greater direct clinical relevance than the ROC curve
- The *no information* classifier always achieves PPV equal to the *base rate, or prevalence* (why?)
- PPV as well as the area under this curve (average precision) must be interpreted relative to prevalence



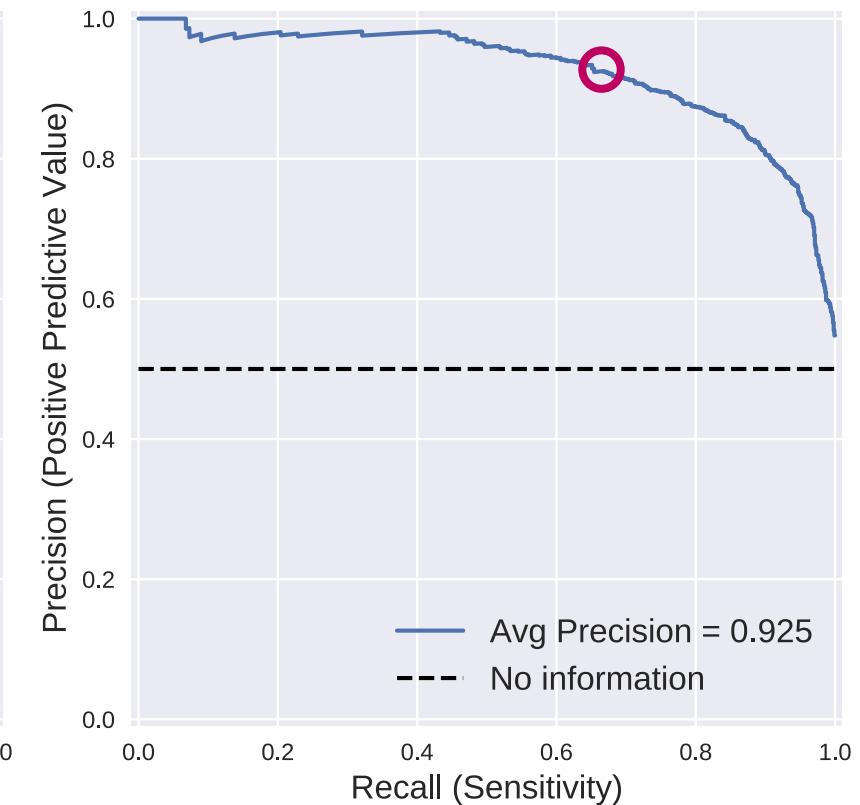
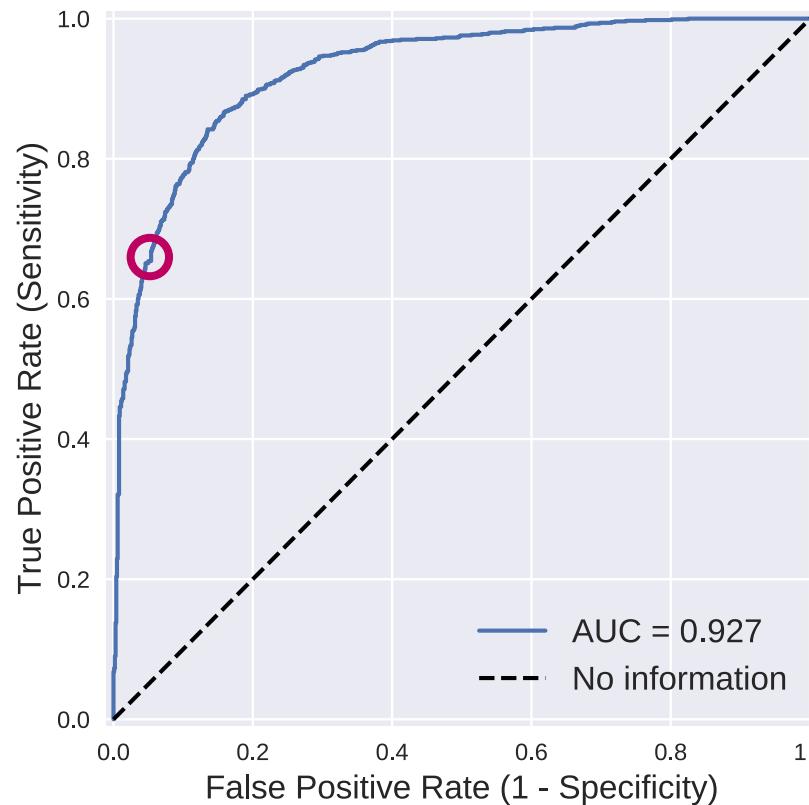
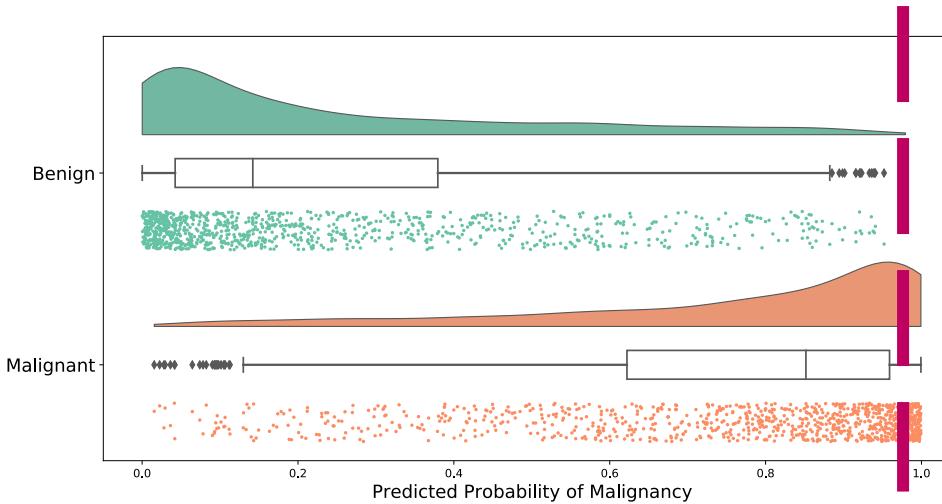
# Operating Point:

*high sensitivity*



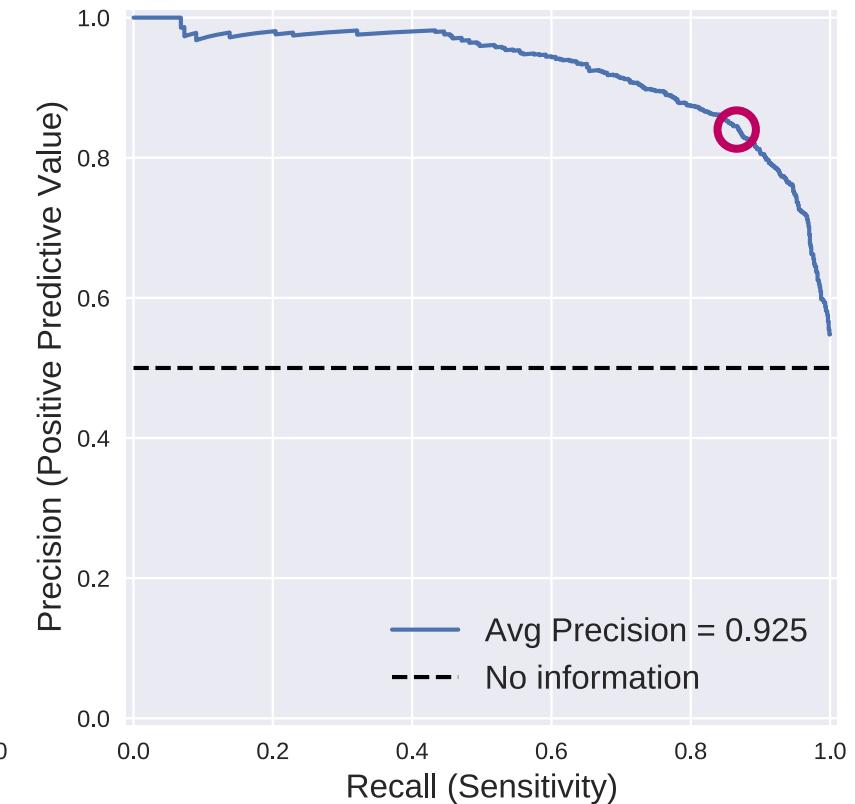
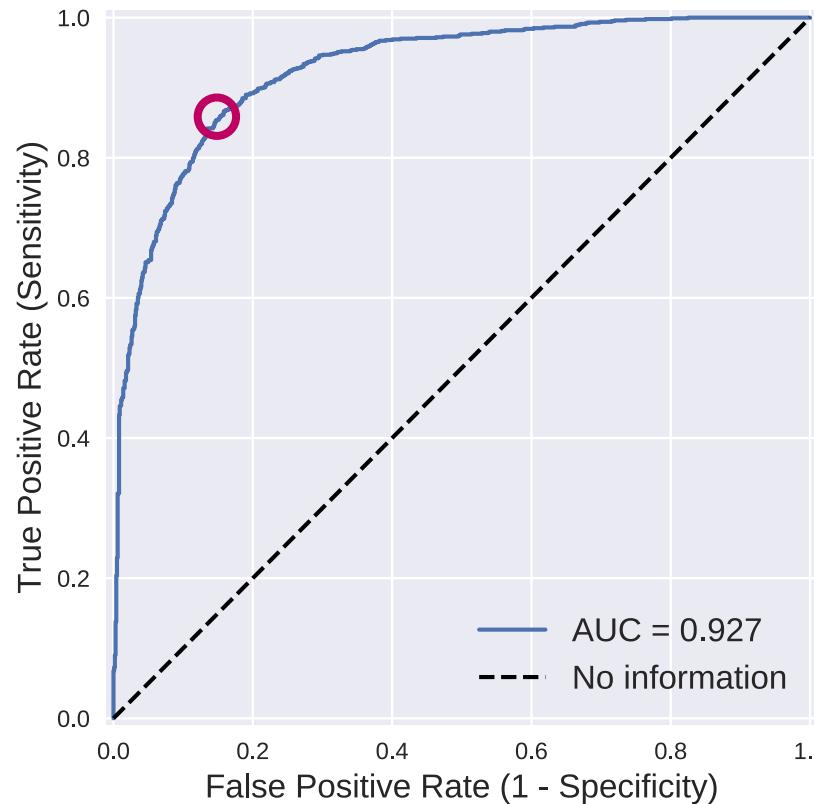
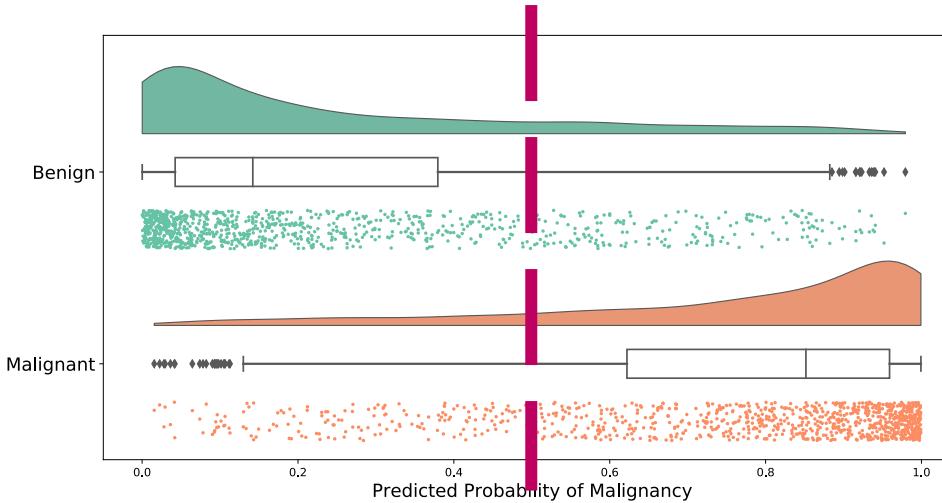
# Operating Point:

*high specificity*



# Operating Point:

*balanced*



# Summary

- It is critical to understand performance measures in order to critically evaluate models and put them to clinical/healthcare use.
- To contextualize performance, we often compare models to a *no information* model whose predictions are random.
- However, *good* performance depends on existing alternative approaches, both tech- and non-tech-based, and the clinical scenario.
- Which measure is most important also depends on the clinical scenario.

# Confusion Matrix for Multi-Class Problems

If time allows.

# Multi-class problems: “Confusion Matrix”

		Predicted Label				
		1	2	3	4	5
True Label	1	True Positive	False Positive	False Positive	False Positive	False Positive
	2	False Negative	True Positive	False Positive	False Positive	False Positive
	3	False Negative	False Negative	True Positive	False Positive	False Positive
	4	False Negative	False Negative	False Negative	True Positive	False Positive
	5	False Negative	False Negative	False Negative	False Negative	True Positive

# Multi-class problems: Binary for Label 1

		Predicted Label				
		0	1	2	3	4
True Label	0	True	False	False	False	False
	1	False	True	False	False	False
	2	False	False	True	False	False
	3	False	False	False	True	False
	4	False	False	False	False	True
	5	False	False	False	False	False

# Multi-class problems: Binary for Label 2

		Predicted Label				
		0	1	2	3	4
True Label	0	Yellow	Blue	Blue	Blue	Blue
	1	Blue	Yellow	Blue	Blue	Blue
2	Blue	Blue	Yellow	Blue	Blue	Blue
3	Blue	Blue	Blue	Yellow	Blue	Blue
4	Blue	Blue	Blue	Blue	Yellow	Blue

There are many more, of course, but classification metrics go a long way.

- Regression
  - Mean squared error (MSE)
  - Mean absolute error (MAE)
  - $R^2$
- Survival Analysis (i.e. failure time)
  - Concordance index
  - MSE, MAE
  - Brier Score
  - $AUC_t$