Applied **Machine Learning**: Predicting **7-year Prostate Cancer** Survivability

Executive Summary

- We can develop many different models model that predict 7-year survival for prostate cancer patients. An **xgBoost** model with a **manual screening step** has the **highest test accuracy**.
- Challenges include:
 - Dataset **dirtiness** many missing values.
 - **Intentional obfuscation** due to many recorded variables not having a clear definition.
- Final model at-a-glance:
 - Gradient-boosted random forest ensemble model + manual screening rule
 - Text parsing + dummy variables + low variance filter + normalization + derivative features
 - Quick Performance Metrics compared to naive null model:
 - Accuracy: **acc** = **0.6875** (+34.47% lift)
 - F1 Measure: **F1** = **0.7187** (**+26.15%** lift)

Research Mission

- To develop a model that:
 - Combines factors to predict **7-year survivability** of prostate cancer patients.
 - Optimizing for accuracy variable importance not explicitly needed.
- Pitfalls:
 - Data points are **intentionally obfuscated**, which might make human intervention to remove spurious variables more difficult.

Dataset Introduction

- Source: **proprietary** collected data from Enova.
- Collection Methods:
 - Unknown. Proprietary data.
- Shape:
 - Rows: **15385**
 - Columns: **33**
- Target Variable shape: binary (1,0).
- Goal: to **predict** patients' **survivability** for diagnosed **Prostate Cancer**.

Dealing with **NAs**

pos Definition	varname	NA	rate.rest	rate.NAs	NA_method	NA_method_justification
18 Size of primary tumor 6 months after diagnosis, in mm	tumor_6_months	0.6540	0.4862	0.4798	remove col	too much missing.
21 Level of prostate-specific antigen in blood 6 months after diagnosis, in ng/mL	psa_6_months	0.6179	0.4859	0.4796	remove col	too much missing.
12 count of family members who have been diagnosed with prostate cancer	family_history	0.1034	0.4810	0.4905	force to 0	predominant subgroup + maybe nothing to record? = 0
13 count of brothers and fathers of the patient who have been diagnosed with prostate cancer	first_degree_histor	0.1034	0.4810	0.4905	force to 0	predominant subgroup + maybe nothing to record? = 0
14 flag indicating whether the patient has ever been diagnosed with any cancer previously	previous_cancer	0.1034	0.4810	0.4905	force to 0	predominant subgroup + maybe nothing to record? = 0
15 flag indicating whether the patient describes himself as a smoker	smoker	0.1034	0.4810	0.4905	force to 0	predominant subgroup + maybe nothing to record? = 0
23 How many times the patient reports drinking tea per week	tea	0.1034	0.4810	0.4905	remove col	not used in modeling
20 Level of prostate-specific antigen in blood at time of diagnosis, in ng/mL	psa_diagnosis	0.0909	0.4816	0.4856	remove	cannot make meaningful prediction without diagnosis info
10 Height of patient at time of diagnosis	height	0.0894	0.4831	0.4704	remove	no default value, cannot impute.
11 Weight of patient at time of diagnosis	weight	0.0856	0.4846	0.4539	remove	no default value, cannot impute.
22 Level of prostate-specific antigen in blood 1 year after diagnosis, in ng/mL	psa_1_year	0.0675	0.4820	0.4823	\=6/diag, fallback 0	impute as no change from diagnosis
8 Age of patient at time of diagnosis	age	0.0478	0.4823	0.4765	remove	no default value, cannot impute.
19 Size of primary tumor 1 year after diagnosis, in mm	tumor_1_year	0.0389	0.4818	0.4860	\=6/diag, fallback 0	impute as no change from diagnosis
24 A list of codes indicating the presence of various symptoms. Meaning has been removed.	symptoms	0.0270	0.4800	0.5550	parsed as empty	nothing to record = 0
3 A measurement of how abnormal the cancer cells look compared to normal cells	gleason_score	0.0207	0.4813	0.5140	remove	small proportion + cannot predict w/o diagnosis
17 Size of primary tumor at time of diagnosis, in mm	tumor_diagnosis	0.0197	0.4823	0.4669	remove	small proportion + cannot predict w/o diagnosis
9 Race of patient	race	0.0112	0.4821	0.4740	remove col	not used in modeling
29 brachytherapy used	brch_thrpy	0.0000	0.4820	0.0000		
27 chemotherapy used	chm_thrpy	0.0000	0.4820	0.0000		
28 crypotherapy used	cry_thrpy	0.0000	0.4820	0.0000		
2 the month and year of diagnosis	diagnosis date	0.0000	0.4820	0.0000	remove col	no observable relationship + "00" dates + irregular data shap
26 hormone therapy used	h_thrpy	0.0000	0.4820	0.0000		
1 An identifier used for scoring dataset	id	0.0000	0.4820	0.0000		
6 Describes whether or not the cancer has spread to distant parts of the body	m_score	0.0000	0.4820	0.0000		
31 multiple therapies used in conjunction	multi_thrpy	0.0000	0.4820	0.0000		
5 Describes whether or not the cancer has spread to the lymph nodes	n_score	0.0000	0.4820	0.0000		
30 prostate surgically removed	rad_rem	0.0000	0.4820	0.0000		
25 external beam radiotherapy used	rd_thrpy	0.0000	0.4820	0.0000		A.C. A.L.A. CIL.
16 What side of the prostate the cancer has been found in	side	0.0000	0.4820	0.0000		Atter NA-filtering:
7 Stage of cancer	stage	0.0000	0.4820	0.0000		10000
32 survived 1 year from diagnosis flag	survival_1_year	0.0000	0.4820	0.0000		After NA-filtering: 10606 rows remaining.
4 Describes local extent of prostate tumor	t score	0.0000	0.4820	0.0000		9

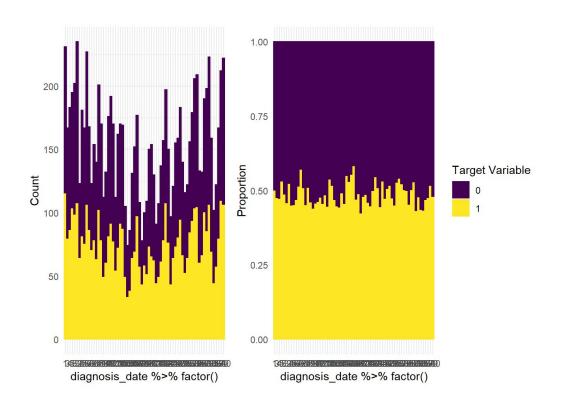
Exploratory Data Analysis #1 - **Discretes**

dableb.				у			isble			У			inlata			у		
variable value n		1	survival	variable	value	n 0		1 survival		variable	value	n	0	1	surviv			
	both	5369	3024	2345	43.68%		0	7123	3978	3145	44.15%	l. 41	0	4894	2431	2463	50.33	
side	left	2045	1154	891	43.57%		1	3020	1759	1261	41.75%	rd_thrpy	1	5712	3568	2144	37.54	
	right	3192	1821	1371	42.95%	family bistons	2	396	220	176	44.44%	la flavor	0	69 59	3972	2987	42.92	
	T1a	672	345	327	48.66%	family_history	3	58	36	22	37.93%	h_thrpy	1	3647	2027	1620	44.42	
	T1b	618	308	310	50.16%		4	7	4	3	42.86%	alama thamas	0	3611	1836	1775	49.16	
	T1c	660	335	325	49.24%		5	2	2	0	0.00%	chm_thrpy	1	69 95	4163	2832	40.49	
	T2a	888	444	444	50.00%		0	8771	4914	3857	43.97%	2000 at 2000	0	8054	4681	3373	41.88	
	T2b	876	441	435	49.66%		1	1672	988	684	40.91%	cry_thrpy	1	2552	1318	1234	48.35	
t_score	T2c	867	464	403	46.48%	first_degree_history	2	143	84	59	41.26%	hards thems.	0	8003	4631	3372	42.13	
	T3a	1075	646	429	39.91%		3	16	10	6	37.50%	brch_thrpy	1	2603	1368	1235	47.45	
	T3b	1105	647	458	41.45%	previous_cancer	4	4	3	1	25.00%	rad_rem 0	0	8744	4980	3764	43.05	
	T3c	1013	602	411	40.57%		0	9960	5624	4336	43.53%		1	1862	1019	843	45.27	
	T4	2832	1767	1065	37.61%		1	646	375	271	41.95%	multi throv	0	2322	1169	1153	49.66	
	N0	6643	3240	3403	51.23%	Landon	0	10079	5717	4362	43.28%	multi_thrpy	1	8284	4830	3454	41.69	
n_score	N1	2895	2191	704	24.32%	smoker	1	527	282	245	46.49%	survival_1_year	0	1107	1107	0	0.00	
	NX	1068	568	500	46.82%		0	765	426	339	44.31%		1	9499	4892	4607	48.50	
	MO	9783	5315	4468	45.67%		1	1837	1025	812	44.20%							
	M1a	297	248	49	16.50%		2	2475	1389	1086	43.88%							
m_score	M1b	184	156	28	15.22%		3	2039	1157	882	43.26%							
	M1c	342	280	62	18.13%		4	1334	772	562	42.13%							
	1	374	127	247	66.04%	4	5	611	354	257	42.06%							
	IIA	1426	597	829	58.13%	tea	6	295	177	118	40.00%							
stage	IIB	2409	1324	1085	45.04%		7	116	64	52	44.83%							
	III	1790	834	956	53.41%		8	37	21	16	43.24%							
	IV	4607	3117	1490	32.34%		9	17	12	5	29.41%							
	1	624	388	236			10	2	1		50.00%							
	2	1560	872	688			12	1	1	0	0.00%							
race	3	427	231	196	45.90%													
	4	7887	4449	3438														

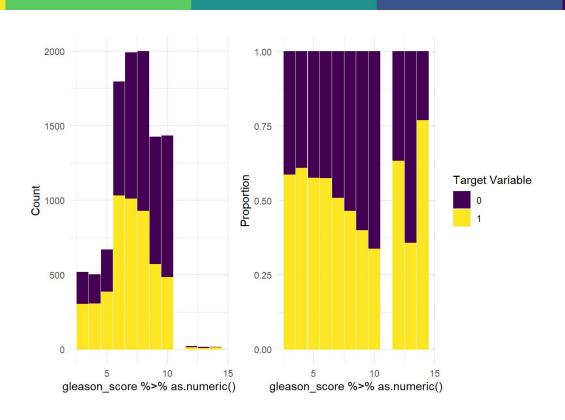
Exploratory Data Analysis #2 - **Symptoms**

variable	volue	D	у		ounival	variable	volue		у		oundival
variable	value	n	0	1	survival	variable	value	n	0	1	survival
overetors 001	0	10150	5163	4987	49.13%	aymentana 1101	0	4067	2083	1984	48.78%
symptom_O01	1	208	167	41	19.71%	symptom_U01	1	6291	3247	3044	48.39%
aumentary 000	0	10233	5228	5005	48.91%	overetore 1102	0	4694	2413	2281	48.59%
symptom_O08	1	125	102	23	18.40%	symptom_U02	1	5 664	2917	2747	48.50%
aumentary 000	0	10260	5247	5013	48.86%	symptom_U03	0	6777	3490	3287	48.50%
symptom_O09	1	98	83	15	15.31%		1	3581	1840	1741	48.62%
aumentary 010	0	10282	5274	5008	48.71%	symptom_U05	0	9339	4712	4627	<mark>49.</mark> 54%
symptom_O10	1	76	56	20	2 6.32%		1	1019	618	401	<mark>39</mark> .35%
aumantana O11	0	8783	4517	4266	48.57%	avmentana 1106	0	8243	4250	3993	<mark>48.</mark> 44%
symptom_O11	1	1575	813	762	48.38%	symptom_U06	1	2115	1080	1035	<mark>48.</mark> 94%
aumantana DO4	0	10012	5062	4950	49.44%	aumantana CO4	0	7813	4032	3781	48.39%
symptom_P01	1	346	268	78	22.54%	symptom_S04	1	2545	1298	1247	<mark>49.</mark> 00%
overstone DO2	0	10133	5151	4982	49.17%	aumentana CO7	0	6229	3251	2978	47.81%
symptom_P02	1	225	179	46	20.44%	symptom_S07	1	4129	2079	2050	49.65%
overstom DO2	0	10287	5270	5017	48.77%	symptom \$10	0	9795	4974	4821	<mark>49.</mark> 22%
symptom_P03	1	71	60	11	15.49%	symptom_S10	1	563	356	207	<mark>36</mark> .77%

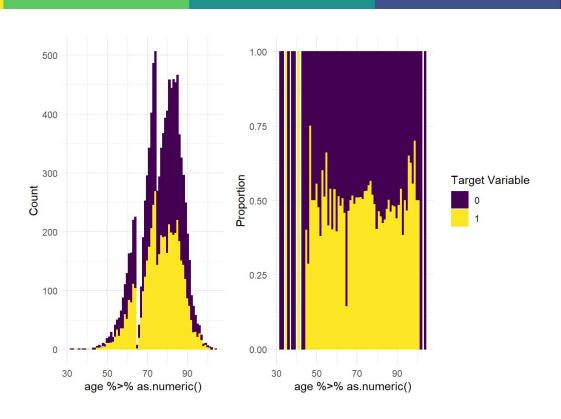
Exploratory Data Analysis #3 - **Numerics**



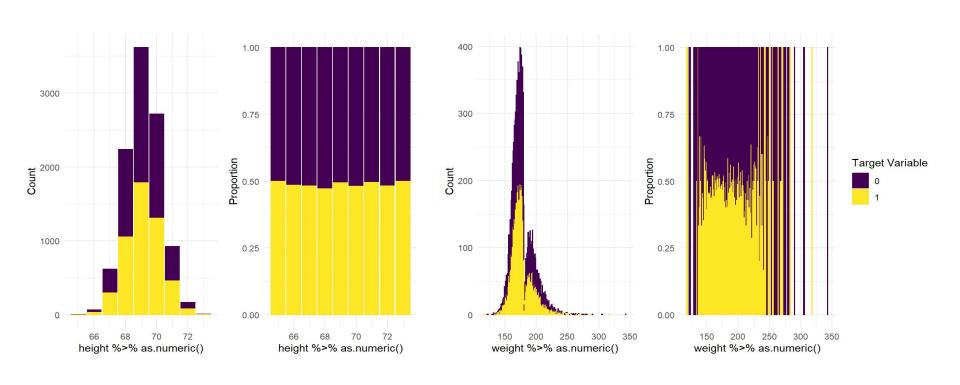
Exploratory Data Analysis #4 - **Numerics**



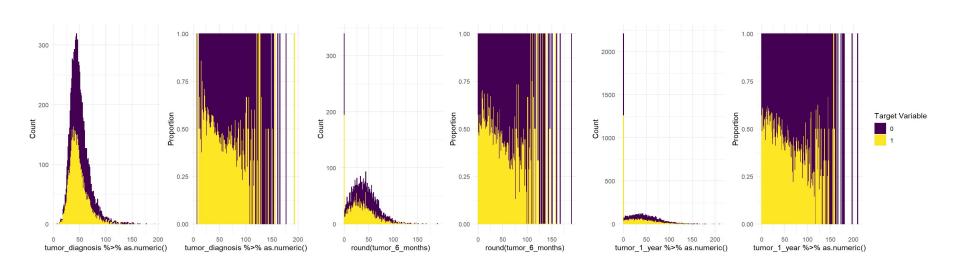
Exploratory Data Analysis #5 - **Numerics**



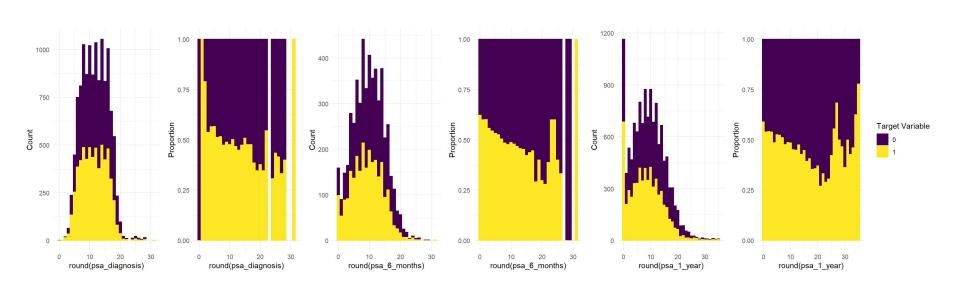
Exploratory Data Analysis #6 - **Numerics**



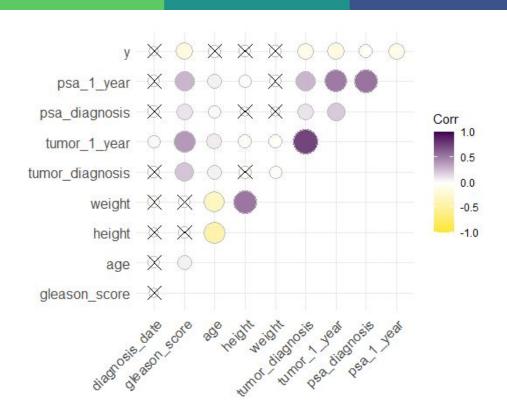
Exploratory Data Analysis #7 - **Numerics**



Exploratory Data Analysis #8 - Numerics



Exploratory Data Analysis #9 - Numerics CorrMap



Train-Test **Splitting**

- Split proportion:
 - 80% training
 - 20% testing

Phase 1 Modeling: Basic Models

- Processing steps:
 - Tokenizing symptom string
 - Deriving Term-Frequency from tokens
 - Set Reference levels for Dummy variables
 - Create **Dummy** variables
 - Filtering for **no-variance** predictors

Basic Models #1: Naive **Sampling** (no features)

- y_{hat} is predicted completely blindly based on distribution of training y=(0,1).

```
blind <- function(dframe){
   dframe %>%
     mutate(
      y_hat = sample(c("0","1"), replace = T, prob = probs.train, size = nrow(dframe))
   )
}
```

model	accuracy	f_meas	lift
Naive Sampling	0.5113101	0.5695309	0.0000000

Basic Models #2: Manual Model

```
manual1 <- function(dframe) {</pre>
  dframe %>%
    mutate(
      y hat = case when(
        # 1-vr Dead Rule
        survival 1 year == 0 ~ 0,
        # Stage Rule
        stage %in% c("IIB", "III", "IV") ~ 0.
        # Metastatisation Rule
        m score != "M0" ~ 0,
        n score != "NO" ~ 0,
        # Symptoms Rule
          symptoms o01 == 1 \sim 0,
          _{\text{symptoms}} oo8 = 1 \sim 0,
          _{\text{symptoms}} == 1 \sim 0,
          symptoms o10 == 1 \sim 0,
          _symptoms_p01 == 1 \sim 0,
          _symptoms_p02 == 1 ~ 0,
          symptoms p03^{\cdot} == 1 \sim 0,
          symptoms s10^{\circ} == 1 \sim 0,
        TRUE ~ 1
```

- Manual model with few rules, and no decision tree. Simple flag-setting.

model	accuracy	f_meas	lift
Manual Rules	0.6140434	0.7310345	0.2009217

Basic Models #3: Naive Bayes

- Default, untuned Naive Bayes model.

model	accuracy	f_meas	lift
Naive Sampling	0.5113101	0.5695309	0.0000000
Naive Bayes	0.5730443	0.7224265	0.1207373
Manual Rules	0.6140434	0.7310345	0.2009217

Basic Models #4: base GLM LogReg

- Default, untuned GLM Logistic Regression Classifier.

term	odds_ratio	p.value	sig	_			
n_score_N1	0.44	0.0000000	***				
gleason_score	0.89	0.0000000	***	model	accuracy	f meas	lift
tumor_1_year	0.99	0.0000000	***	Section 19 and 1			
tf_symptoms_u05	0.65	0.0000002	***	Naive Sampling	0.5113101	0.5695309	0.0000000
rd_thrpy	0.75	0.0000134	***	Naive Bayes Manual Rules	0.5730443 0.6140434	0.7224265 0.7310345	0.1207373 0.2009217
tf_symptoms_s10	0.64	0.0001020	***	base GLM	0.6743638	0.7075751	0.3188940
rad_rem	0.79	0.0012385	**				
weight	0.99	0.0012616	**				
brch_thrpy	0.81	0.0020365	**				

Phase 2 Modeling: Intermediate Models

- Additional Feature Engineering:
 - Calculating BMI & binning into weight classes (under/normal /overweight/obese)
 - Calculating PSA & tumor size change: 1-year delta
 - **Centering & scaling** all

 NUMERICS (for regularised GLM family only xgBoost & tree family does not benefit from scaling)

model	accuracy	f_meas	lift
Naive Sampling	0.5113101	0.5695309	0.0000000
Naive Bayes	0.5730443	0.7224265	0.1207373
Manual Rules	0.6140434	0.7310345	0.2009217
base GLM	0.6743638	0.7075751	0.3188940

Phase 2 Modeling: Intermediate Models

- Cross-validation split: **4-fold CV**, 75/25. CV on d.train only.
- Parameter Tuning:
 - Parameters:
 - **Elastic-net GLM**: 6x6 standard sampling grid
 - penalty: how much regularisation?
 - mixture : what type of regularisation?
 - **xgBoost**: length 30 Latin hypercube sampling grid
 - learn rate : learning rate.
 - loss_reduction : required loss reduction for further node-split
 - min_n: minimum data points at a node for further node-split
 - mtry: proportion of predictors randomly sampled at split
 - sample size : proportion of data used in fitting
 - tree_depth : max tree depth.

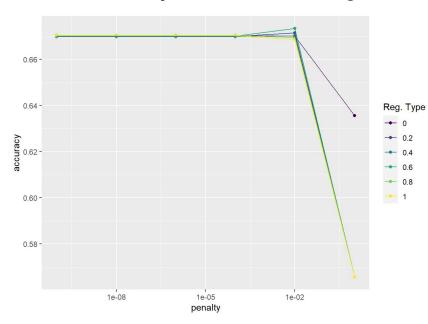
Intermediate Models #1: base GLM w/ FE

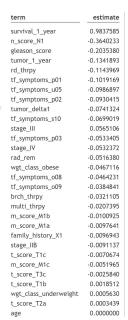
- **Unpenalised GLM** logistic classifier model. Uses feature-engineered dataset.

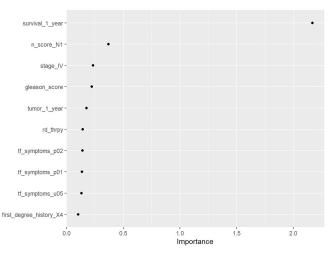
term	estimate	p.value	sig				
n_score_N1	-0.3688596	0.0000000	***				
gleason_score	-0.2207874	0.0000000	***	model	accuracy	f_meas	lift
tumor_1_year	-0.3135726	0.0000000	***	Naive Sampling	0.5113101	0.5695309	0.0000000
tf_symptoms_u05	-0.1323507	0.0000002	***	Naive Bayes	0.5730443	0.7224265	0.1207373
rd_thrpy	-0.1435097	0.0000122	***	Manual Rules base GLM	0.6140434 0.6743638	0.7310345 0.7075751	0.2009217 0.3188940
tf_symptoms_s10	-0.1026216	0.0000985	***	engineered GLM	0.6757776	0.7084746	0.3216590
wgt_class_obese	-0.0924944	0.0004049	***				
rad_rem	-0.0921186	0.0012108	**				
brch_thrpy	-0.0868727	0.0024341	**				

Intermediate Models #2: elastic-net GLM

- Tuned, **penalised GLM** logistic classifier model. Uses feature-engineered dataset.



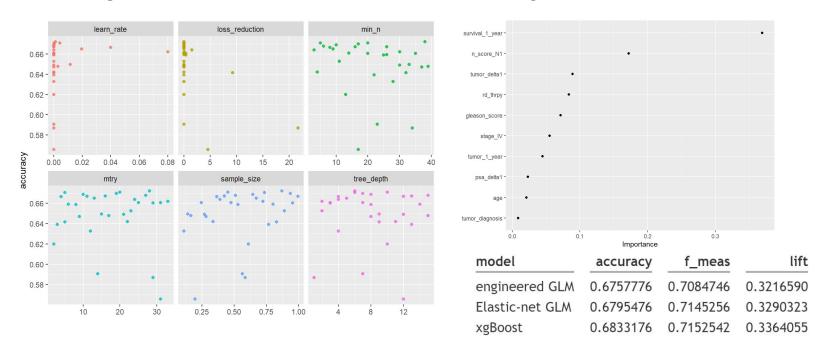




model	accuracy	f_meas	lift	
engineered GLM	0.6757776	0.7084746	0.3216590	
Elastic-net GLM	0.6795476	0.7145256	0.3290323	

Intermediate Models #3: xgBoost

- Tuned, **xgBoost** classifier model. Uses feature-engineered dataset.



Intermediate Models: Model Performances

model	accuracy	f_meas	lift
Naive Sampling	0.5113101	0.5695309	0.0000000
Naive Bayes	0.5730443	0.7224265	0.1207373
Manual Rules	0.6140434	0.7310345	0.2009217
base GLM	0.6743638	0.7075751	0.3188940
engineered GLM	0.6757776	0.7084746	0.3216590
Elastic-net GLM	0.6795476	0.7145256	0.3290323
xgBoost	0.6833176	0.7152542	0.3364055

Phase 3 Modeling: the Human Touch

- Recall that one of the variables is survival_1_year. This is a perfect predictor for y=0 (not y=1), but some models might not be able to converge on this rule, which could be a way to squeeze out some more performance from the model.
- Additionally, we can add more screening rules from the manual model earlier.
- Thus, we stack this screening step on a model. Experimental code:

```
# Stacking a check for survival_1_year on top of any fitted model
manual_stack <- function(dframe, modelfit){

df_man <- dframe[dframe$survival_1_year == min(dframe$survival_1_year),]%>%
    mutate(.pred_class = 0)

df_alg <- dframe[dframe$survival_1_year != min(dframe$survival_1_year),]
    df_alg <- df_alg %>% cbind(predict(modelfit, new_data = df_alg))

result <- rbind(df_man, df_alg)
    return(result)
}</pre>
```

Phase 3 Modeling: the Human Touch

Final Metrics:

model	accuracy	f_meas	lift
Naive Sampling	0.5113101	0.5695309	0.0000000
Naive Bayes	0.5730443	0.7224265	0.1207373
Manual Rules	0.6140434	0.7310345	0.2009217
base GLM	0.6743638	0.7075751	0.3188940
engineered GLM	0.6757776	0.7084746	0.3216590
Elastic-net GLM	0.6795476	0.7145256	0.3290323
Manual + Elastic-net	0.6795476	0.7145256	0.3290323
xgBoost	0.6833176	0.7152542	0.3364055
Manual + xgBoost	0.6875589	0.7187102	0.3447005