

Predicting Malignancy from Mammography Findings and Surgical Biopsies

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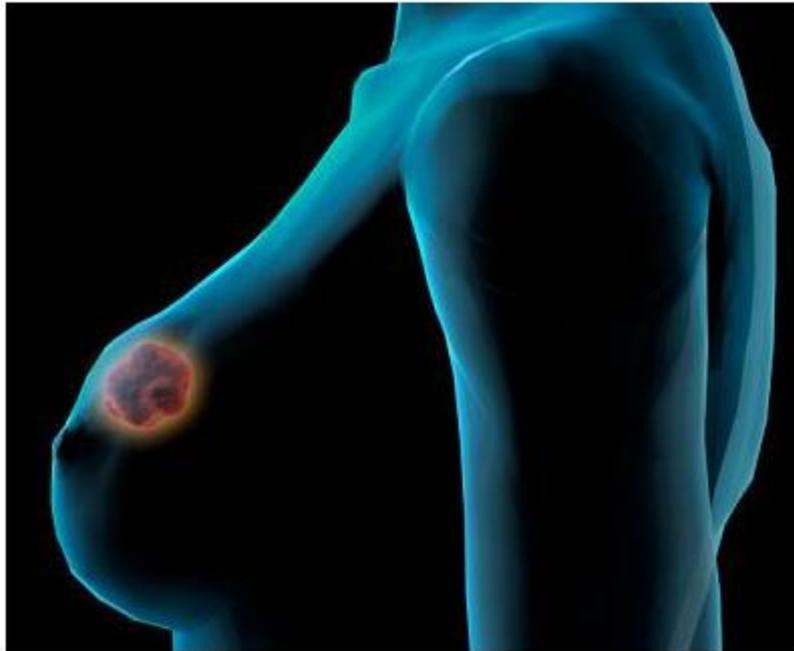
Outline

- Breast Cancer
- Objectives
- Data
- Methodology
- Results and Analysis
- *MammoClass* (online application)
- Conclusions and Future Work

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Breast Cancer



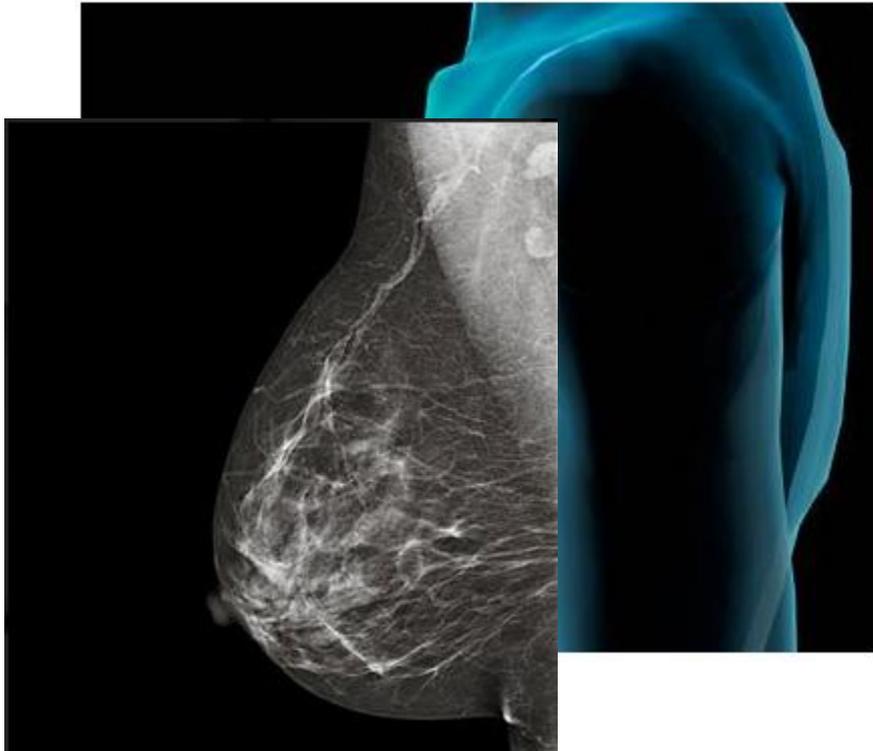
- USA:
 - 1 woman dies of breast cancer every 13 minutes
 - In 2011:
 - 230.480 invasive cancers
 - 39.520 ($\approx 17\%$) expected to die

Source: *U. S. Breast Cancer Statistics* –
October 2011

- Portugal:
 - Per year:
 - 4500 new cases
 - 1500 deaths (33%)

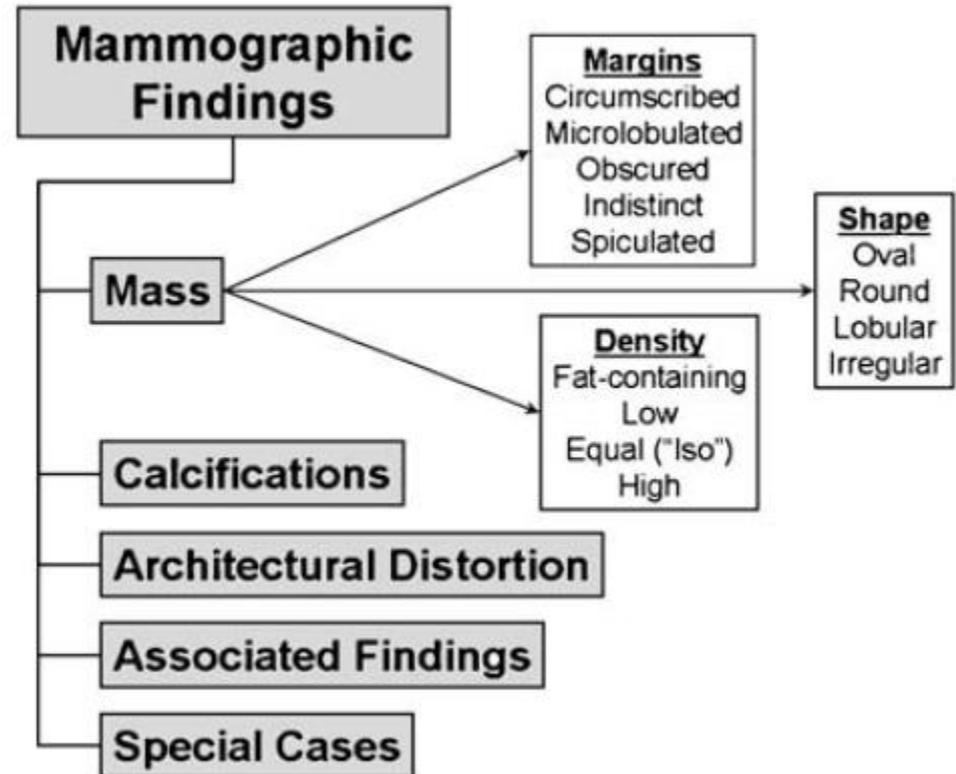
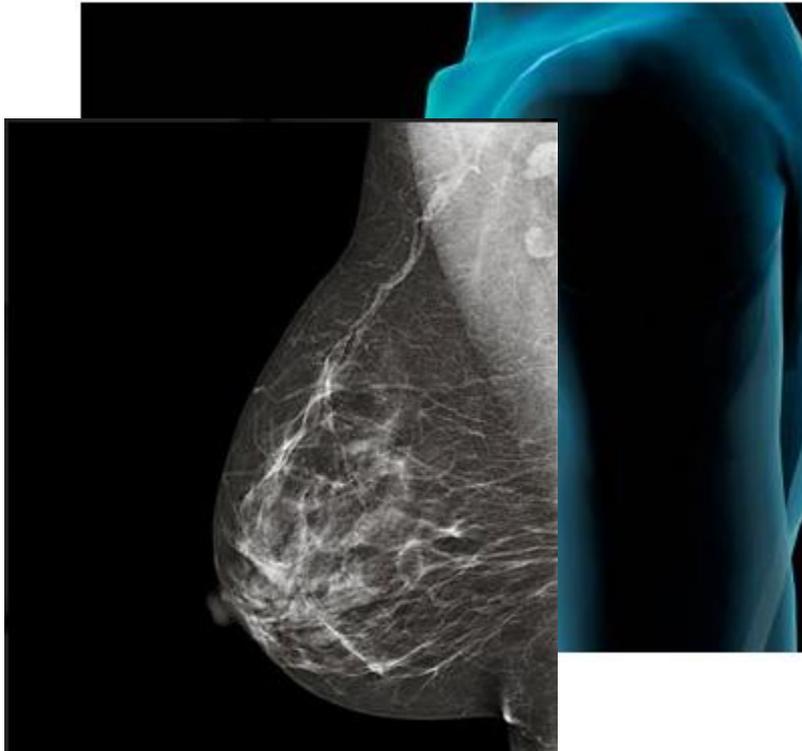
Source: *Liga Portuguesa Contra o Cancro* –
November 2011

Breast Screening Programs



- Reduction of death rate in 30%
- **Mammography:**
The cheapest and most efficient method to detect cancer in a preclinical stage

Mammography



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in *Studying the relevance of Breast Imaging Features* – HEALTHINF 2011

Objectives



- Build classifiers capable of predicting **mass density** and **malignancy** from a reduced set of mammography findings



- Reduce the number of unnecessary biopsies

Outline

- Breast Cancer
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Data



- Source:
 - Ryan Woods (M.D.)
 - Elizabeth Burnside (M.D.)



- 348 cases
- Each case refers to a breast nodule **retrospectively** classified according to BI-RADS[®] system
- From mammographies results
- Collected between October 2005 and December 2007

Attributes

age_at_mammo

CLOCKFACE_LOCATION_OR_REGION

MASS_SHAPE

MASS_MARGINS

SIDE

DEPTH

MASS_MARGINS_worst

QUADRANT_LOCATION_def

SIZE

OVERALL_BREAST_COMPOSITION

Density_num

retro_density

outcome_num

Masses classification

Prospective

- **Classification** of feature **mass density** for **180** cases **just by one radiologist**:
 - low density;
 - iso-dense;
 - high density;
- **Brief** and superficial medical **report** (at the time of imaging);
- **Classification under stress.**



mass density

density_num

Retrospective

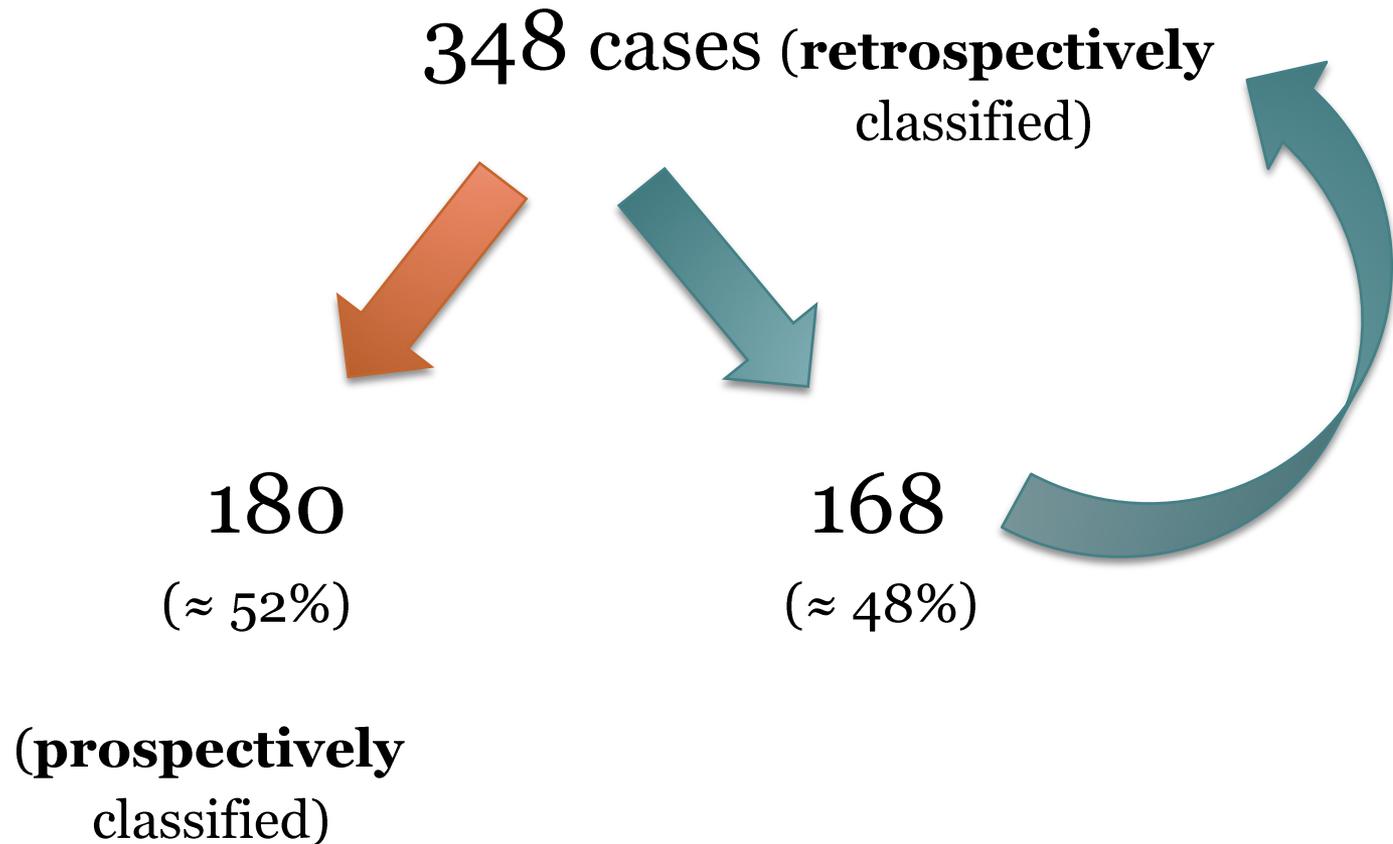
- **Classification** by a **group of experienced physicians** that **re-assess** all exams (**348**);
- **Review of mass density classification** made by radiologist (prospective study);
- **Classification without stress**;
- **Reference standard** for **mass density.**



mass density

retro_density

Masses classification



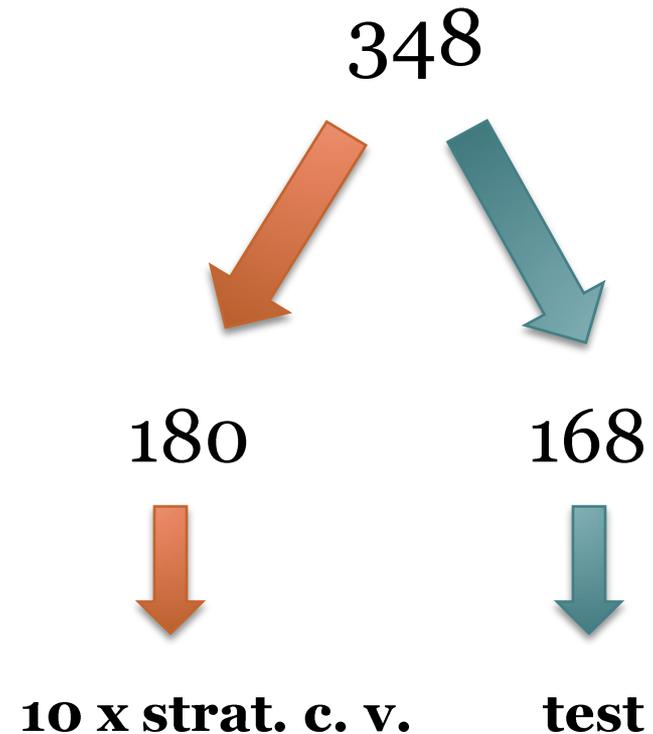
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Methodology



- WEKA
- Paired Corrected T-Tester
 - **Significance level: 0.05**



Methodology - Algorithms applied

- ZeroR (baseline classifier)
- OneR
- DTNB
- PART

rules

- J48
- DecisionStump
- RandomForest
- SimpleCart
- NBTree

trees

- NaiveBayes
- BayesNet (TAN)

bayes

- SMO } *functions*

functions

internal parameter variation

Methodology - Experiments

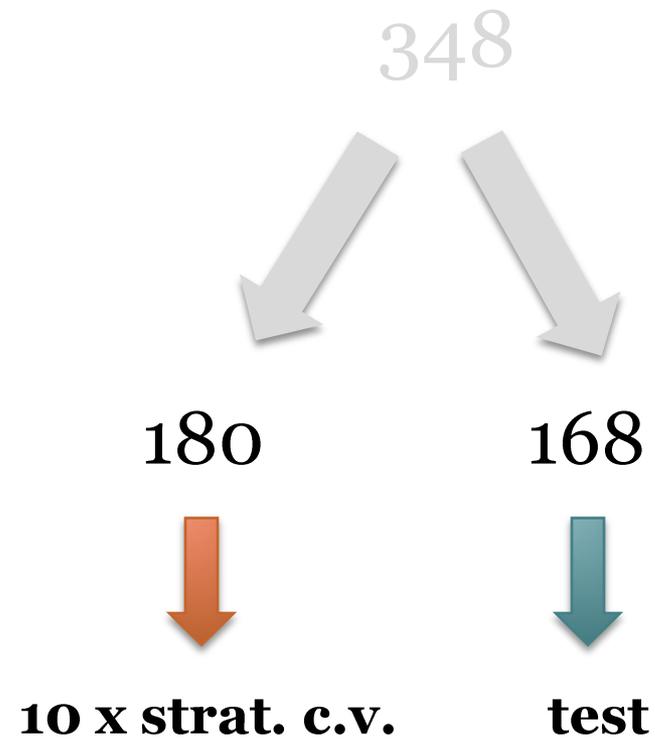
10 x stratified. c. v.

180

- E_1 – Predicting malignancy with *retro_density*
- E_2 – Predicting malignancy with *density_num*
- E_3 – Predicting malignancy without mass density

- E_4 – Predicting *retro_density*
- E_5 – Predicting *density_num*

Results



Results - Experiments

10 x stratified. c. v.

180

Exp.	Algorithm	CCI	K	F	AUROC
E1	SMO	85.6±7.3	0.69±0.16	0.80±0.11	0.84±0.08
E1	DTNB	81.6±8.2	0.60±0.18	0.74±0.13	0.88±0.07
E1	NaiveBayes	81.3±9.5	0.61±0.20	0.76±0.12	0.88±0.08
E1	J48	80.7±9.3	0.59±0.20	0.75±0.13	0.79±0.11
E2	SMO	83.9±7.7	0.66±0.17	0.78±0.11	0.82±0.08
E2	NaiveBayes	80.3±9.3	0.59±0.19	0.75±0.12	0.87±0.09
E2	DTNB	79.8±9.5	0.56±0.21	0.72±0.15	0.86±0.09
E2	J48	75.4±9.5	0.47±0.21	0.65±0.15	0.73±0.12
E3	SMO	83.8±7.7	0.65±0.17	0.78±0.11	0.82±0.09
E3	J48	76.3±9.9	0.49±0.22	0.67±0.15	0.76±0.13
E3	NaiveBayes	76.2±9.9	0.51±0.20	0.71±0.13	0.85±0.09
E3	DTNB	75.7±9.0	0.48±0.19	0.67±0.13	0.81±0.10
E4	SMO	81.3±8.2	0.52±0.21	0.64±0.17	0.75±0.11
E4	J48	74.4±8.8	0.32±0.24	0.47±0.21	0.67±0.15
E4	DTNB	73.5±10.0	0.34±0.24	0.51±0.19	0.76±0.12
E4	NaiveBayes	72.8±9.9	0.37±0.23	0.56±0.18	0.77±0.11
E5	NaiveBayes	67.2±12.1	0.33±0.25	0.62±0.15	0.72±0.14
E5	SMO	66.8±10.7	0.31±0.22	0.55±0.16	0.65±0.11
E5	J48	63.6±10.1	0.26±0.21	0.56±0.15	0.62±0.13
E5	DTNB	62.1±11.9	0.22±0.24	0.54±0.16	0.64±0.14

Results - Experiments

Predicting density

180

Results - Experiments

10 x stratified. c. v.

- E_4 – Predicting *retro_density*

SVM's

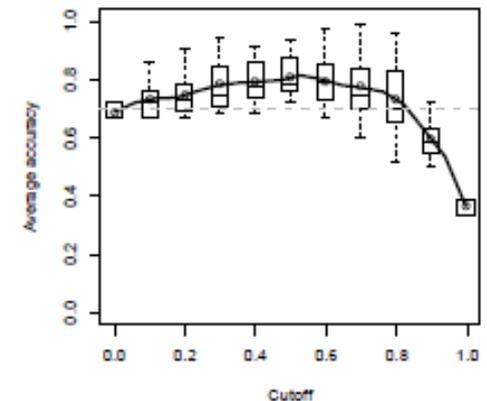
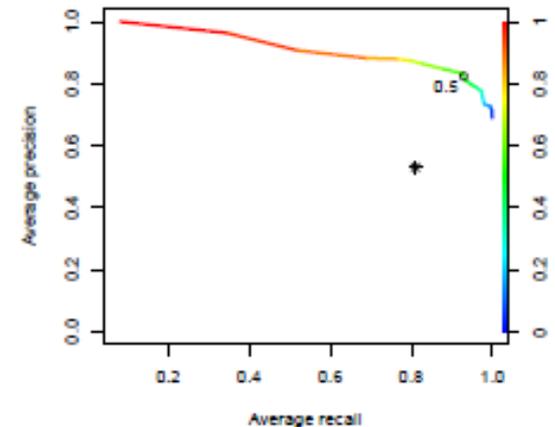
CCI: 81.3% (+/-8.2)

K: 0.52 (+/- 0.21)

F: 0.64 (+/- 0.17)

Radiologist's accuracy = 70 %

Our classifier \approx 81 %



Results - Experiments

TEST

- **E₆** – Predicting *retro_density*
(model E₄ applied)

SVM's

CCI: 84.5%

K: 0.46

F: 0.91

180

SVM's

CCI: 81.3% (+/-8.2)

K: 0.52 (+/- 0.21)

F: 0.64 (+/- 0.17)

Results - Experiments

Predicting malignancy

180

Results - Experiments

10 x stratified. c. v.

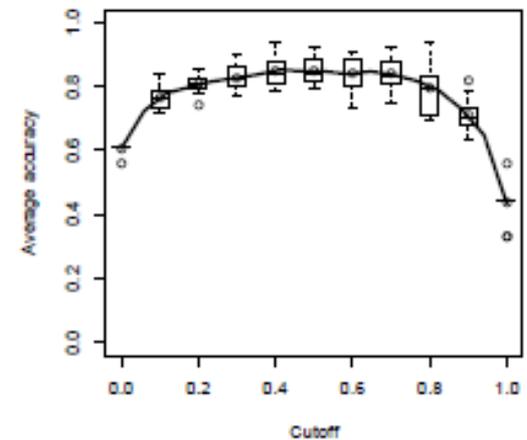
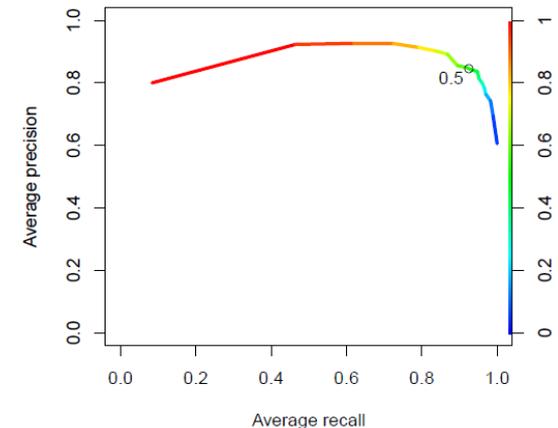
- E_1 – Predicting malignancy with *retro_density*

SVM's

CCI: 85.6% (+/-7.3)

K: 0.69 (+/- 0.16)

F: 0.80 (+/- 0.11)



168

Results - Experiments

TEST

180

SVM's

CCI: 85.6% (+/-7.3)

K: 0.69 (+/- 0.16)

F: 0.80 (+/- 0.11)

- E_8 – Predicting malignancy with *retro_density*
(model E_1 applied)

SVM's

CCI: 81.0%

K: 0.50

F: 0.87

 with **real** values
of **retro_density**

SVM's

CCI: 78.0%

K: 0.45

F: 0.85

 with **predicted**
values of
retro_density
by classifier E_6

MammoClass

- Online application freely available at:
 - <http://cracs.fc.up.pt/mammoclass/>

MammoClass

Classification of a mammogram based in a reduced set of mammography findings

To obtain a prediction in terms of malignancy for a certain mass is only necessary to provide the values of the findings, annotated through the Breast Imaging Reporting and Data System (BIRADS), in the form bellow. It is also possible to get a prediction of the attribute *mass density* in case this feature is not known.

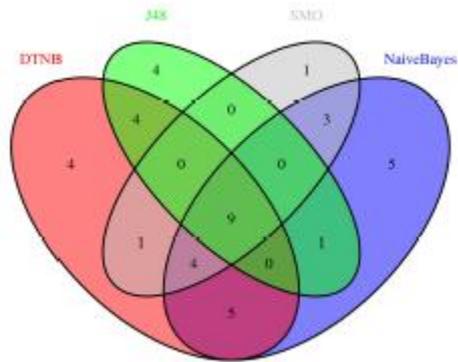
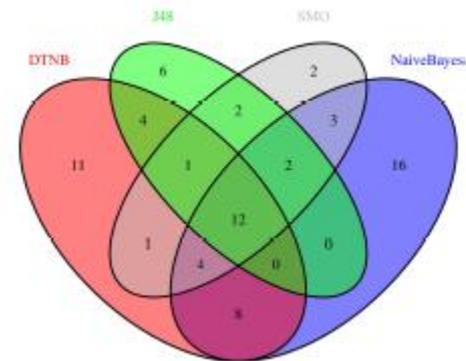
The output will indicate the probability of a certain mass being benign or malignant. In the latter case it is suggested that the patient should perform a biopsy. The probabilities are computed using machine learning models built as described in:

- P.Ferreira, N. A. Fonseca, I. Dutra, R. Woods, and E. Burnside, ***Predicting Malignancy from Mammography Findings and Surgical Biopsies***

Enter Data

Patient's age
 Mass size
 Breast Composition
 Mass shape
 Mass clockface location
 Mass margins (1)
 Mass margins (2)

Misclassified Instances


 E_1

 E_4

Conclusions and Future Work

- a) **Automatic classification** of a mammography can reach **equal or better results** than the classification performed by a **radiologist**;
- b) Machine learning **classifiers** can **predict mass density** with **higher quality** than the one obtained by radiologists
 - a) our classifier can **predict malignancy** in the absence of mass density, since we can **fill up** this **attribute** using our **mass density predictor**.

Conclusions and Future Work

- a) Apply other machine learning techniques based on statistical relational learning;
- b) Investigate how other features can affect malignancy or are related to the other attributes;
- c) Study why the **parameter variation** on **WEKA algorithms** has a strong **impact** on the **performance** of **classifiers**;
- d) Investigate with the radiologist why some **instances** are **consistently misclassified** by all algorithms.

Thank you!



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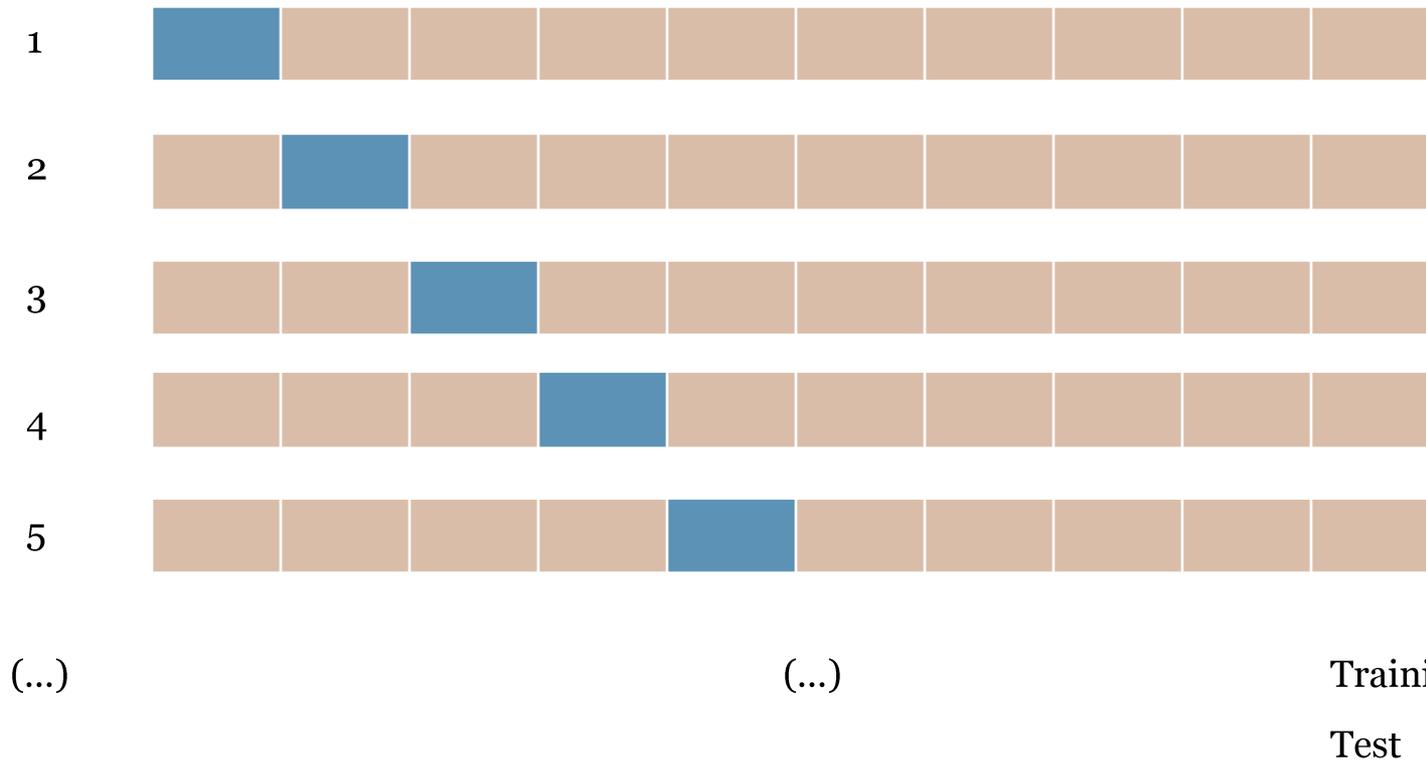
<http://cracs.fc.up.pt>

Appendices

Methodology

10-fold stratified cross-validation

Iteration



Data distribution

- 348

348	<i>retro_density</i>		Total
<i>outcome_num</i>	<i>high</i>	<i>iso</i>	
<i>malignant</i>	59 (70.2%)	59 (22.3%)	118 (33.9%)
<i>benign</i>	25 (29.8%)	205 (77.7%)	230 (66.1%)
Total	84 (24.1%)	264 (75.9%)	

Data distribution

- 180

180	<i>retro_density</i>		Total
<i>outcome_num</i>	<i>high</i>	<i>iso</i>	
<i>malignant</i>	42 (75.0%)	29 (23.4%)	71 (39.4%)
<i>benign</i>	14 (25.0%)	95 (76.6%)	109 (60.6%)
Total	56 (31.1%)	124 (68.9%)	

180	<i>density_num</i>		Total
<i>outcome_num</i>	<i>high</i>	<i>iso</i>	
<i>malignant</i>	51 (63.0%)	20 (20.2%)	71 (39.4%)
<i>benign</i>	30 (37.0%)	79 (79.8%)	109 (60.6%)
Total	81 (45.0%)	99 (55.0%)	

Data distribution

- 168

168	<i>retro_density</i>		Total
<i>outcome_num</i>	<i>high</i>	<i>iso</i>	
<i>malignant</i>	17 (60.7%)	30 (21.4%)	47 (28.0%)
<i>benign</i>	11 (39.3%)	110 (78.6%)	121 (72.0%)
Total	28 (16.7%)	140 (83.3%)	

WEKA algorithms used

CLASSIFIERS' PERFORMANCE FOR EACH TASK. VALUES NOT IN BOLD ARE STATISTICALLY SIGNIFICANTLY WORSE THAN THE CLASSIFIER WITH HIGHEST ACCURACY (USING PAIRED T-TEST WITH $\alpha = 0.05$).

Exp.	Algorithm	CCI	K	F	AUROC
E1	SMO	85.6 \pm 7.3	0.69 \pm 0.16	0.80 \pm 0.11	0.84 \pm 0.08
E1	DTNB	81.6 \pm 8.2	0.60 \pm 0.18	0.74 \pm 0.13	0.88 \pm 0.07
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E4	NaiveBayes	72.8 \pm 9.9	0.37 \pm 0.23	0.56 \pm 0.18	0.77 \pm 0.11
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E5	DTNB	62.1 \pm 11.9	0.22 \pm 0.24	0.54 \pm 0.16	0.64 \pm 0.14

Parameter variation in WEKA algorithms

(E_1) Predicting *outcome_num* with *retro_density*,

(E_2) Predicting *outcome_num* with *density_num*,

(E_3) Predicting *outcome_num* without mass density,

(E_4) Predicting *retro_density*,

(E_5) Predicting *density_num*,

Best parameter selection

SMO '-C 0.05 -N1 PK -E 1.0'

SMO '-C 0.05 -N2 PK -E 1.0'

naïve Bayes default

Parameters Selection:

SMO:

-C (complexity parameter)

-N (filterType)

0 - Normalize training data

1 - Standardize training data

2 - No normalization/standardization

PK (PolyKernel)

-E (exponent value)

debug	False
displayModelInOldFormat	False
useKernelEstimator	False
useSupervisedDiscretization	False