

Deep Learning methods for Quality Check Algorithms in Mammography

Research Internship (April-October 2019)

Author: Baptiste Doyen

Supervisors: Pierre Fillard (CSO)

Yaroslav Nikulin (*Sr. Research Scientist*)

Therapixel - CentraleSupélec/MVA

6th September 2019 - Paris Saclay campus

Context: Breast Cancer, AI & Quality check

The most diagnosed and lethal cancer for women with **1/4** of total cancers and **627K** deaths worldwide^[9]

- Recurrent screenings
- CAD assistance
- New generation with AI
- AI require both High Volume and High Quality Data

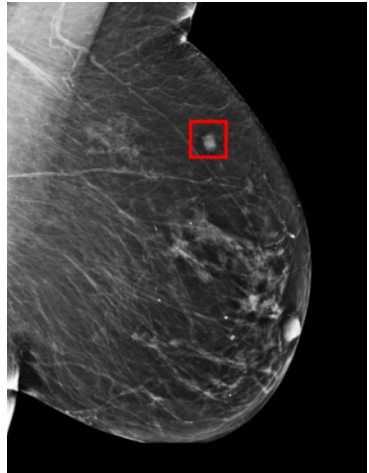


Figure 1: A malignant lesion on a Mammography

Quality Check

A **crucial step** for Cancer Detection algorithms.
Traditionnally performed **manually** but today **automatized**

Two main steps:

- **Pre-Quality Check:** separate screening images from non-screening images
(global features of the image)
- **Geometry Check:** ensure that breast geometry on the mammography is correct
(more local and constrained features of the image)

Pre-Quality Check

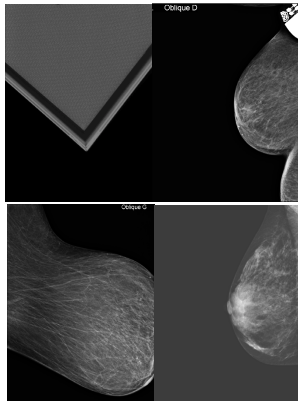


Figure 2: Non-Screening

VS.

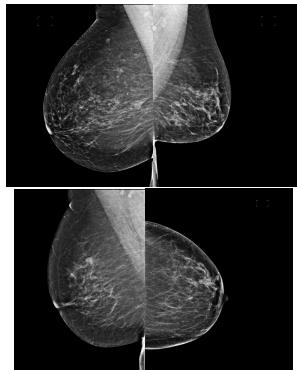


Figure 3: Screening

Geometry Check

The goal is to extract feature points from the image and check if they comply with set of rules S_{geom}

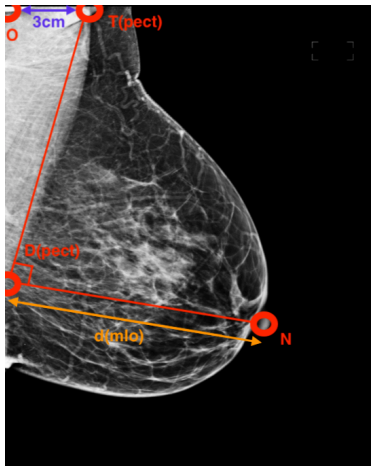


Figure 4: MLO view

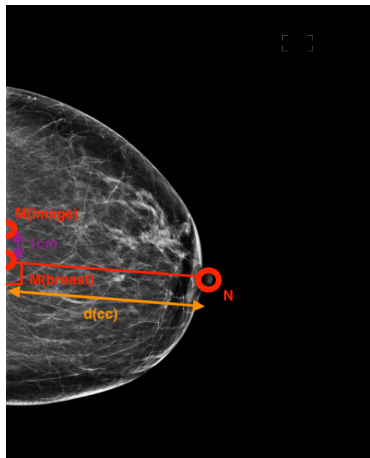
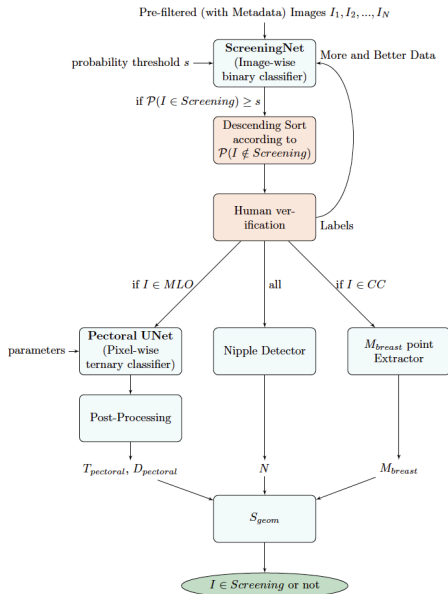


Figure 5: CC view

Decision pipeline



Decision pipeline (bis)

- In blue : automatic processes
- In orange : human assistance to improve these processes (~ anomaly detection)
- My work was mainly focused on **ScreeningNet**, **Pectoral UNet** and **Post-Processing**

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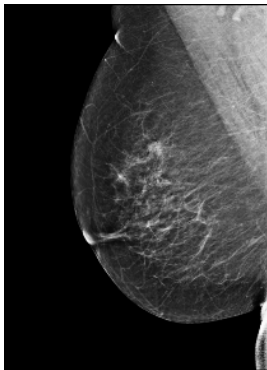
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Mammography Segmentation

Visual example:



$$x_0 \xrightarrow{f^*} y_0$$



Problem definition:

- $r, c \in \mathbb{N}$
- $x \in \mathbb{R}^{r \times c}$ a mammography image
- $y \in \{0, 1\}^{r \times c \times 3}$ a Ground-Truth (GT) tensor
- ℓ a segmentation loss function measuring discrepancy between masks
- $\mathcal{F} \subset (\{0, 1\}^{r \times c \times 3})^{\mathbb{R}^{r \times c}}$ a function space

We look for:

$$f^* = \underset{f \in \mathcal{F}}{\operatorname{argmin}} \ell(f(x), y) \quad (1)$$

Non Deep Learning methods

Test of two classical approaches:

1) Frontier estimation with
Image gradient and
Continuous line search

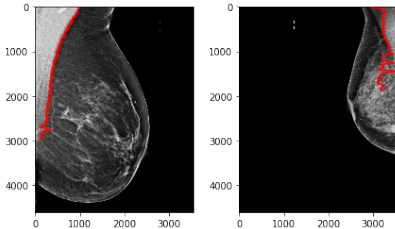


Figure 6: Success and Failure

2) FloodFill^[1] algorithm

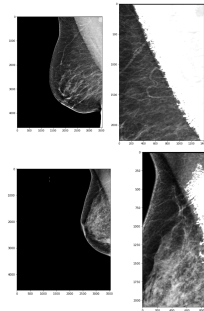


Figure 7: Success and Failure

⇒ good results only on easy cases

Deep Learning methods

Method based on **learning**:

we assume $\mathcal{F} = \mathcal{F}_\Theta$ (with $\Theta \in \mathbb{R}^N$ and $N \gg 1$) and we intend to learn a set of parameters Θ^* by optimizing ℓ on Θ space.

More details:

- A similar approach has been developed in 2017 by Alejandro Rodriguez-Ruiz et al.^[4] (reference paper)
- **Advantages:**
 - extract more advanced image *features* (e.g. texture)
 - only few labelled data
- **Drawbacks:**
 - overfit happens and prevent *reproducibility*
 - domain *transferability*

UNet Network: f_{Θ}

A **symmetric** and **fully-convolutional** architecture made of:

- **Contracting network:** *what is there?*
- **Expanding network:** *where is it?*
- short and long **Skip connections**

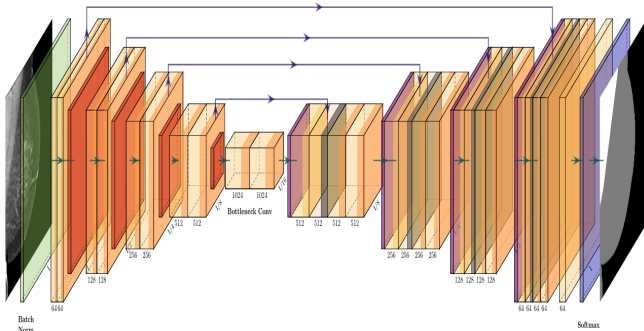


Figure 8: end-to-end architecture from Image pixels to GT labels

Loss function: ℓ

Inspired from UNet paper^[6] and also from Isensee et al.^[2]

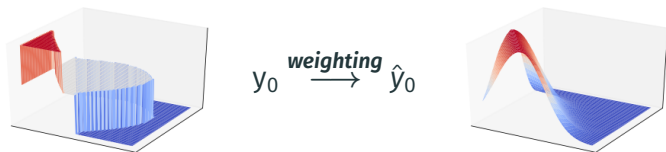
$$\ell(f_{\Theta}(x), y) = (1 - \lambda) \underbrace{\left(- \sum_{c \in C} \sum_{i,j} w_x(i,j) y_{i,j,c} \log(f_{\Theta}(x)_{i,j,c}) \right)}_{\mathcal{L}_{CE}} + \lambda \underbrace{\left(- \frac{2}{|C|} \sum_{c \in C} \frac{\sum_{i,j} f_{\Theta}(x)_{i,j,c} y_{i,j,c}}{\sum_{i,j} f_{\Theta}(x)_{i,j,c} + \sum_{i,j} y_{i,j,c}} \right)}_{\mathcal{L}_{Dice}} \quad (2)$$

with: $[f_{\Theta}(x)_{i,j,c}]_{i,j,c}$ output tensor, $[y_{i,j,c}]_{i,j,c}$ GT tensor, C the set of classes, λ a ponderation parameter and $[w_x(i,j)]_{i,j}$ the weight-map associated to x .

Weight-map: w_x

- Adaptation of Cross-Entropy (CE) for **borders zone**
- Equivalent to a **'new' GT distribution**:

$$\hat{y}_{i,j} = w_x(i,j) \bullet y_{i,j} \quad (3)$$



- **Stochastic Gradient Descent**

- **Gradient**: Backpropagation^[5] to get $\nabla_{\Theta} \ell(f_{\Theta}(x), y)$
- **Optimizer**: Adam^[3]

- **Metrics** to assess final performances:

- **Dice** Median :

$$D([x_{i,j,c}], [y_{i,j,c}]) = \frac{2 \times |\{(i,j), x_{i,j,c=pect}=1\} \cap \{(i,j), y_{i,j,c=pect}=1\}|}{|\{(i,j), x_{i,j,c=pect}=1\}| + |\{(i,j), y_{i,j,c=pect}=1\}|}$$

- Inter-Quartile Range (**IQR**)

Data & Pre-Processing

- **169** images labelled manually with software *labelme*^[8] (121 training/15 validation/33 test)
- 2 main **constraints** on Data Augmentation:
 - preserve **minimal pectoral presence**
 - preserve **spatial relative arrangement**



Figure 9: From left to right *Missing pectoral, non-natural border and mirroring*

Which model configuration performs the best ?

- Impossible to test all hyper-parameter combinations!

But:

- Accurate enough *a priori* values
- Not full-independence, $batchSize \Rightarrow N_{epochs}, N_{improve}, d$
- 2 regimes ($batchSize \in \{6, 10\}$) and learnings from one transferred to the second
- We first fixed a *benchmark model* for each regime and iterations with *small incremental changes* one at a time.

Results

- Best model configuration:

$d = 1\%$ | $batchSize = 10$ | BN=Yes | L. ReLU | Inv=No |
 $\lambda_{loss} = 0.5$ | Depth=5 | flips+elastic | $N_{init-filters} = 64$

- Final results:

Model(s)	Manufacturer	Result
<i>Single</i> best	Hologic	0.9647 (0.0243)
{ <i>Single</i> + <i>PP</i> } best	Hologic	0.9683 (0.0224)
<i>Ensemble</i> best	Hologic	0.9654 (0.0213)
{ <i>Ensemble</i> + <i>PP</i> } best	Hologic	0.9654 (0.0201)
Ref. ^[4] best on DBT	Siemens	0.977 (0.0170)
Ref. ^[4] best on DM	Siemens	0.974 (0.0170)
Ref. ^[4] best on DM	Hologic	0.947 (0.0559)

ScreeningNet

- Binary **classification** problem to ease Screening process
- **Model used:** modified VGG16^[7]-like network with $\sim 10K$ images
- **Global result:** $AUC = 0.8259$ (validation of the approach)
- **More in details:** the aim is to minimize FPR (False Positive Rate). For instance, $FPR = 5\% \implies TPR = 18\%$ (still low)

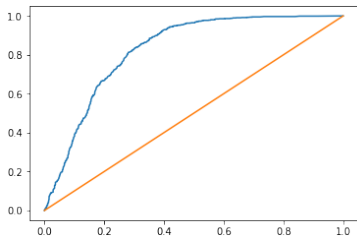


Figure 10: ROC Curve with $AUC = 0.8259$

Conclusion & Future work

- **Quality Check:**
requires both **non-DL & DL** stand-alone expert modules
- **On Mammography Segmentation:**
results **similar** to the reference paper^[4].
At first sight: same approach but some changes were necessary to achieve **reproducibility**.
»Next steps: address domain **transferability** issue
- **On ScreeningNet:**
general approach **validated** but *TPR* is still too low.
»Next steps: More data and multi-class approach.

New framework for Mammography Segmentation

- **Motivations:** noisy/leaky/uncomplete results with UNet-only. **Post-Processing** is here to address this issue but can be improved.
A learnt and thus **optimized** post-processing would be better.
- **SegAN^[10] framework:** **GAN** approach with UNet as Mask Generator and a deep multi-scale concatenated network (*critic*) as Discriminator.

Main innovation: one single loss to train both networks

$$\ell_{SegAN} = l_{MAE} \left[f_{\Theta_{Critic}} \left([x_{i,j}] \bullet [f_{\Theta_{UNet}}(x_{i,j})]_{c=pectoral} \right), \right. \\ \left. f_{\Theta_{Critic}} \left([x_{i,j}] \bullet [y_{i,j,c}]_{c=pectoral} \right) \right] \quad (4)$$

New framework for Mammography Segmentation

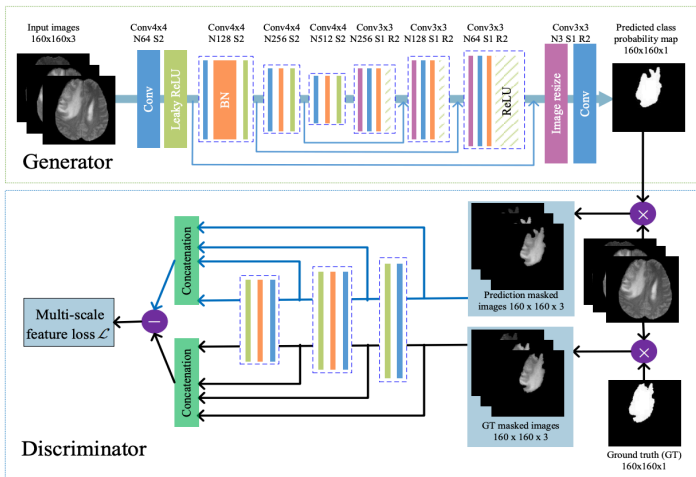


Figure 11: SegAN architecture. Image Credit to Xue Yuan et al.^[10]

Thanks for your attention! :)

References

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- [2] Isensee, F. et al. (2018). nnU-Net: Self-adapting Framework for U-Net-Based Medical Image Segmentation. *ArXiv e-prints*.
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- [6] Ronneberger, O. et al. (2015). U-Net: Convolutional Networks for Biomedical Image Segmentation. *ArXiv e-prints*.
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- [8] Wada, K. (2016). labelme: Image Polygonal Annotation with Python. <https://github.com/wkentaro/labelme>.
- [9] WHO (2018). <https://www.who.int/cancer/PRGlobocanFinal.pdf>.