## CHAPTER 3

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# Biomedical Texture Operators and Aggregation Functions

A Methodological Review and User's Guide

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## Abstract

This chapter reviews most popular texture analysis approaches under novel comparison axes that are specific to biomedical imaging. A concise checklist is proposed as a user guide to assess the relevance of each approach for a particular medical or biological task in hand. We revealed that few approaches are regrouping most of the desirable properties for achieving optimal performance. In particular, moving frames texture representations based on learned steerable Riesz operators showed to enable data-specific and rigid-transformationinvariant characterization of local directional patterns, the latter being a fundamental property of biomedical textures. Potential limitations of having recourse to data augmentation and transfer learning for deep convolutional neural networks and dictionary learning approaches to palliate the lack of large annotated training collections in biomedical imaging are mentioned. We conclude by summarizing the strengths and limitations of current approaches,

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providing insights on key aspects required to build the next generation of biomedical texture analysis approaches.

Keywords: Biomedical texture analysis, convolutional neural networks, deep learning, dictionary learning, Gabor filters, gray-level co-occurrence matrices, gray-level run-length matrices, gray-level size-zone matrices, local binary patterns, steerable wavelets.

## 1. Introduction

When starting from scratch to enable quantitative image analysis in a biomedical research project or a clinical setting, the number of available approaches can be disconcerting. Whereas implementing intensity-based features already raises several challenges in terms of reproducibility and relevance to a particular medical or biological applicative context, finding adequate texture measures requires extensive expertise as well as time-consuming iterative validation processes. Knowing precisely the type of textural information sought and further opting for the appropriate analysis technique is as much challenging as crucial for success. To that end, Section 2 of Chapter 1 proposed a formal definition of Biomedical Texture (BT) from both perceptual and mathematical perspectives. It was suggested that BT are realizations of intricate and non-stationary spatial stochastic processes, and that spatial scales and directions are fundamental properties of BT. The relation between scale and directions was further developed in Section 4.1 of Chapter 2, suggesting that the Local Organization of Image Directions (LOID) is a fundamental property of BT. The LOIDs relate to the texton theory (see Section 2.4 of Chapter 1), where textons are crucial elements of preattentive vision [33]. Section 3.1 of Chapter 1 introduced a general framework of texture analysis methods where any BTA approach can be decomposed into a succession of local texture operators and regional aggregation functions. In addition, Section 3.3 of Chapter 1 and Chapter 7 provided evidence that the type of handcrafted geometric invariances required for BTA are invariances to non-rigid transformations<sup>1</sup>, which considerably differs from general purpose texture analysis traditionally used in computer vision for photographic image analysis. These aspects were further clarified and exemplified in Chapter 2, where a set of comparison dimensions between BTA methods was introduced. In particular, non-exclusive categories of texture operators were presented, including (i) directionally insensitive, (ii) directionally sensitive, (iii) aligned, (iv) Moving Frames (MF) and (v) learned. The discrimination abilities of each category was evaluated, where the destructive effects of integrative aggregation

<sup>1</sup>Optimal operators should be equivariant to translations and locally equivariant to rotations.

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Figure 3.1: Overview and dependence of the BTA approaches reviewed in this chapter. These categories were chosen based on their popularity and diversity.

were demonstrated. In particular, it was found that some groups of operators lose their ability to characterize the LOIDs when integrated over a Region Of Interest (ROI), the latter being required to obtain scalar-valued texture measurements. The ultimate challenging requirement of combining the ability to describe the LOIDs with robustness to rigid transformations was only fulfilled by two similar groups of operators: aligned and MFs.

In this work, we compare most popular BTA approaches under the light of the novel comparison dimensions introduced in Chapter 2. A concise checklist is proposed to assess the relevance of each BTA approach for a particular medical or biological task in hand. The main groups of methods considered are (i) convolutional (Section 2, further information on CNNs can be found in Chapters 4 and 9), (ii) Gray-Level Matrices (GLM, Section 3), (iii) Local Binary Patterns (LBP, Section 4), and (iv) fractals (Section 5, further information can be found in Chapter 5). The main categories of BTA methods are summarized in Fig. 3.1. Our choice to review these categories is based on their popularity and diversity. The strengths and weaknesses of the methods are summarized and discussed in Section 6. The notations used in this chapter are based on Section 2.2 of Chapter 1.

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## 2. Convolutional approaches

A large group of approaches called *convolutional* are based on linear texture operators. In spatial domain, the application of a linear operator  $G_n$  to  $f(x)^2$  at the position  $x_0$  is characterized by the *operator function*  $g_n(x)$  as

$$
\mathcal{G}{f}(x_0) = (g_n * f)(x) = \int_{x \in \mathbb{R}^D} f(x)g_n(x_0 - x) dx.
$$
 (3.1)

The operation  $(g_n * f)(x_0)$  is called a *convolution*: we slide the operator function  $g_n(x)$ to the input texture function  $f(x)$  over all positions  $x_0$ . When the position  $x_0$  is fixed, we remark that the value of  $G(f)(x_0)$  is simply the scalar product

$$
\mathcal{G}_n\{f\}(\mathbf{x}_0) = \langle f(\cdot), g_n(\mathbf{x}_0 - \cdot) \rangle. \tag{3.2}
$$

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A convolution in the spatial domain corresponds to a multiplication in the Fourier domain as

$$
(g_n * f)(x) \xrightarrow{f} \hat{g}_n(\omega) \hat{f}(\omega).
$$

Convolution operators are the ones for which the response map  $h_n(x)$  depends linearly on the input texture function  $f(x)$ . The definition of the function  $g_n(x)$  is a priori free of constraints. For the convolution, the spatial support of the operator is exactly the domain on which the operator function  $g_n(x)$  is nonzero. Operators with finite spatial supports are desirable to study local texture properties of non-stationary processes (see Section 3.1 of Chapter 2), hence we shall consider localized operator functions. Also, band-pass filters for which  $\hat{g}_n(\mathbf{0}) = 0$  (*i.e.*, zero gain for the null frequency  $\omega = \mathbf{0}$ corresponding to the average) are able to focus on texture alone and do not include any intensity information. This ensures improved robustness of the texture operator responses to variations in illumination.

As will be exploited thereafter, it is often interesting to define new texture operators based on convolution operators, with the goal of locally aligning the orientation. We develop this framework in the 2D setting. The matrix rotation with angle  $\theta_0$  is noted

$$
\mathbf{R}_{\theta_0} = \begin{pmatrix} \cos \theta_0 & -\sin \theta_0 \\ \sin \theta_0 & \cos \theta_0 \end{pmatrix}.
$$

Consider a convolution operator  $G_n$  with operator function  $g_n(x)$ . For each location  $x_0$ , we set

$$
\theta_{\mathbf{x}_0} = \arg \max_{\theta_0 \in [0, 2\pi)} \langle f(\cdot), g_n(\mathbf{x}_0 - \mathbf{R}_{\theta_0} \cdot) \rangle \tag{3.3}
$$

<sup>2</sup>In this section, except as otherwise stipulated, we will consider continuous operators and functions indexed by the coordinate vector *x*. Discretized versions can be obtained following the notions introduced in the Section 2.2 of Chapter 1.

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Here,  $\theta_{x_0}$  is the angle that maximizes the scalar product between the texture image  $f(x)$  and the rotated version of  $g_n(x)$  around the location  $x_0$ . It is therefore the angle for which the operator function is the most aligned with  $f(x)$  at position  $x<sub>0</sub>$ . We then define

$$
\mathcal{H}_n\{f\}(\mathbf{x}_0) = \langle f(\cdot), g_n(\mathbf{x}_0 - \mathbf{R}_{\theta_{\mathbf{x}_0}} \cdot) \rangle
$$
\n
$$
= \int_{\mathbf{x} \in \mathbb{R}^D} f(\mathbf{x}) g_n(\mathbf{x}_0 - \mathbf{R}_{\theta_{\mathbf{x}_0}} \mathbf{x}) d\mathbf{x}.
$$
\n(3.4)

The response map  $h_n(x) = H_n\{f\}(x)$  is no longer linear in  $f(x)$ , therefore the texture operator  $\mathcal{H}_n$  is not a convolution operator in the sense of Eq. (3.1). We include the angle alignment in this section since it is based on the convolutional framework and characterized by the operator function  $g_n(x)$  according to Eq. (3.4). Convolution operators and local angle alignment together allow to design texture operators equivariant to rotations that can be used to study the local orientation in the texture image.

The most popular convolutional approaches in texture analysis are detailed in the following subsections while distinguishing three main categories of convolutional operator functions (see Fig. 3.1): circularly/spherically symmetric filters (Sec. 2.1), directional filters (Sec. 2.2), and learned filters (Sec. 2.3). It is worth noting that directional and learned filters are non-exclusive categories.

## 2.1. Circularly/spherically symmetric filters

Circularly/spherically symmetric filters are convolutional texture operators with functions that only depend on the radial polar coordinate  $r$  (see Section 2 of Chapter 2):  $g_n(x) = g_n(||x||) = g_n(r)$ . A direct consequence of this is their complete lack of directional sensitivity.

A simple example of such convolutional texture processing is based on Laplacian of Gaussian (LoG) filters<sup>3</sup>. Their handcrafted operator function  $g_{\sigma}(x)$  is a radial second-order derivative of a D-dimensional Gaussian filter as

$$
g_{\sigma}(\mathbf{x}) = -\frac{1}{\pi \sigma^2} \left( 1 - \frac{\|\mathbf{x}\|^2}{2\sigma^2} \right) e^{-\frac{\|\mathbf{x}\|^2}{2\sigma^2}},
$$
(3.5)

where the standard deviation of the Gaussian  $\sigma$  controls the scale of the operator. LoGs are band-pass and circularly/spherically symmetric. It is straightforward to notice in Eq. (3.5) that  $g_{\sigma}(x)$  only depends on the norm of x and is therefore circularly symmetric: we have  $g_{\sigma}(x) = g_{\sigma}(r)$ . Their 2D profiles and response maps obtained with synthetic tumors are depicted in Fig. 1.12 of Chapter 1. LoGs can also be approximated by a difference of two Gaussians<sup>4</sup>. LoGs were mentioned to be important in

<sup>3</sup>They are also called *Mexican hat* filters due their 2D shape.

<sup>&</sup>lt;sup>4</sup>The Difference of Gaussians (DoG) best approximates LoG when the ratio of the two variances of the

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## Table 3.1: Properties LoG filters.



biological visual processing by Marr in [43].

Multi-scale texture measurements  $\eta$  can be obtained by averaging the absolute values or the energies of the response maps (*i.e.*,  $|h_{\sigma_n}(x)|$  or  $h_{\sigma_n}^2(x)$ , respectively) of a series of operators with increasing values of  $\sigma_1 < \sigma_2 < \cdots < \sigma_N$  (see Fig. 1.12 of Chapter 1). They are implemented in 2D in the  $TextRAD<sup>5</sup>$  commercial medical research software [23]. The properties of LoG filters are summarized in Table 3.1.

## 2.2. Directional filters

Several handcrafted approaches were proposed for texture analysis using directional filters. The latter are sensitive to image directions (see Section 4.2 of Chapter 3). The most important and popular methods are discussed in this section, including Gabor wavelets (Section 2.2.1), Maximum Response 8 (MR8, Section 2.2.2), Histogram of Oriented Gradients (HOG, Section 2.2.3), and the Riesz transform (Section 2.2.4).

Gaussians are  $\sigma_1 = \frac{\sigma_2}{\sqrt{2}}$ . <sup>5</sup>http://texrad.com, University of Sussex, Brighton, UK, as of 21 November 2016.

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#### *2.2.1. Gabor wavelets*

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A popular example is the family of Gabor wavelets [36, 52, 29], which consists in a systematic parcellation of the Fourier domain with elliptic Gaussian windows (see Fig. 3.2). The D-dimensional filter function of the Gabor operator  $g_{\rho,\sigma,\theta}(x)$  is the most general function that minimizes the uncertainty principle [50] (see Section 3.1 of Chapter 2). Moreover, it was suggested in [9] that 2D Gabor filters are appropriate models of the transfer function of simple cells in the visual cortex of mammalian brains and thus mimicking the early layers of human visual perception. In the spatial domain, they correspond to Gaussian-windowed oscillatory functions as

$$
g_{\rho,\sigma,\theta}(\mathbf{x}) = \frac{\sigma_1 \sigma_2}{2\pi} e^{-\sigma_1^2 \tilde{x}_1^2 + \sigma_2^2 \tilde{x}_2^2} e^{j2\pi \rho \tilde{x}_1}, \tag{3.6}
$$

where  $\rho$  is the center of the Gaussian window in Fourier,  $\sigma = (\sigma_1, \sigma_2)$  contains the radial and orthoradial standard deviations of the Gaussian window in polar Fourier, and  $\tilde{\mathbf{x}} = (\tilde{x}_1, \tilde{x}_2) = \mathbf{R}_{\theta} \mathbf{x}$  defines the radial and orthoradial elliptic Gaussian axes in space.

Multi-scale and multi-orientation texture measurements  $\eta$  are obtained by averaging the absolute values or the energies of the response maps (*i.e.*,  $|h_{\rho,\sigma,\theta}(x)|$  or  $h_{\rho,\sigma,\theta}^2(x)$ , respectively) for various values of  $\rho$ ,  $\sigma$ , and  $\theta$ . The properties of Gabor wavelets are summarized in Table 3.2.

## *2.2.2. Maximum Response 8 (MR8)*

Another popular example of handcrafted directional filterbank is the MR8 approach [66]. The latter includes a filterbank with a collection of 38 operator functions  $g_n(x)$  (see Fig. 3.3). Two of them are circularly symmetric (one Gaussian and one LoG). Eighteen are directional multi-scale edge detectors based on oriented first-order Gaussian derivatives. Another eighteen are directional multi-scale ridge detectors based on oriented second-order Gaussian derivatives. However, the approach yields a total of eight response maps. The first two come from the convolution of the image with the circularly symmetric Gaussian and LoG filters. The remaining six are computed as follows. For each directional detector types (*i.e.*, first- and second-order Gaussian derivatives), only one detector per scale is kept. Among the six different filter orientations per scale, the detector kept at the position  $x_0$  is the one that maximizes the detection in the sense of Eq. (3.3), where  $\theta_0$  is coarsely discretized. This results in six texture operators  $G_{n=1,\dots,6}$  that are "aligned" at every position and achieve approximated local rotation-invariance. This process is illustrated in Fig. 3.3.

The aggregation function consists in pixel-wise clustering of the maximum filter responses to create a texton dictionary (see Section 2.4 of Chapter 1). Class-wise models can then be created as texton occurrence histograms, which can be used to compare texture instances by measuring distances between them  $(e.g., Euclidean, \chi^2)$ . The vector of texture measurements  $\eta$  contains the bin values of the texton occurrences. The

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## Table 3.2: Properties of Gabor wavelets.



properties of the MR8 approach are summarized in Table 3.3.

## *2.2.3. Histogram of Oriented Gradients (HOG)*

An efficient gradient-based MF representation is the local image descriptor used in the Scale Invariant Feature Transform (SIFT) approach [40] called Histogram of Oriented Gradients (HOG). The texture function  $f(x)$  is first filtered with circularly symmetric multi-scale DoG filters *g*σ*<sup>i</sup>* (*x*) (see Section 2.1) using a dyadic scale progression (*i.e.*,  $\sigma_{i+1} = 2\sigma_i$ , which yields a collection of response maps  $h_{\sigma_i}(x)$ . At a fixed scale *i*, corresponding to the standard deviation  $\sigma_i = \sigma$ , the gradient orientation map  $h^{\theta}_{\sigma}(\mathbf{x})$  is computed from the response map  $h_{\sigma}(x)$  as

$$
h_{\sigma}^{\theta}(\boldsymbol{x}) = \arctan\left(\frac{\nabla_{x_2} \{h_{\sigma}\}(\boldsymbol{x})}{\nabla_{x_1} \{h_{\sigma}\}(\boldsymbol{x})}\right),\tag{3.7}
$$

 $06$ The texton dictionary is derived from the data, but the operators are handcrafted.

 $07$ Clustering results in averaging within local homogeneous regions in the feature space.

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Figure 3.2: One isolated scale  $\rho$  of Gabor wavelets with 4 orientations  $\theta$  and a spectral dispersion of  $\sigma = (\sigma_1, \sigma_2)$ . Note that the spatial angular polar coordinate  $\theta$  is equivalent to the Fourier angular coordinate  $\vartheta$ .

where  $\nabla_{x_1} \{h_{\sigma}(x) \}$  and  $\nabla_{x_2} \{h_{\sigma}(x) \}$  yield gradients maps. Eq. (3.7) provides local angle values maximizing the gradient magnitude in the spirit of Eq. (3.3). For a fixed orientation  $\theta$ , we set  $u(\theta) = \mathbf{R}_{\theta}(1, 0) = (\cos \theta, \sin \theta)$  and  $\nabla_{u(\theta)} = \cos \theta \cdot \nabla_{x_1} + \sin \theta \cdot \nabla_{x_2}$ the oriented derivative with direction  $u(\theta)$ . We define the oriented response map as  $h_{\sigma,\theta}$ { $x_0$ } =  $\nabla$ <sub>*u*( $\theta$ </sub>){ $h_{\sigma}$ }( $x_0$ ). When using discretized image functions, operators and response maps  $h_{\sigma}(\xi)$ , the latter can be estimated using pixel differences as

$$
\nabla_{x_1} \{ h_{\sigma} \}(\xi) = h_{\sigma}(\xi_1 + \Delta \xi_1, \xi_2) - h_{\sigma}(\xi_1 - \Delta \xi_1, \xi_2),
$$
  
\n
$$
\nabla_{x_2} \{ h_{\sigma} \}(\xi) = h_{\sigma}(\xi_1, \xi_2 + \Delta \xi_2) - h_{\sigma}(\xi_1, \xi_2 - \Delta \xi_2).
$$
\n(3.8)

For each position  $x_0$ , the dominant gradient direction  $\theta_{x_0}$  is obtained by maximizing Eq. (3.7). To define the HOG operators, we align and sample the orientation  $\theta$  =  $\theta_q - \theta_{x_0} = \frac{2\pi}{q} - \theta_x$  with  $q = 1, ..., 8$ . The final collection of HOG operators  $\mathcal{G}_{\sigma,q}$  is given at location  $x_0$  by

$$
\mathcal{G}_{\sigma,q}\{f\}(\boldsymbol{x}_0)=h_{\sigma,q}(\boldsymbol{x}_0).
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## Table 3.3: MR8 properties.



The value of  $h_{\sigma,q}(x_0)$  can be efficiently evaluated using the steerability of the gradient operator [21] as

$$
h_{\sigma,q}(\mathbf{x}_0) = \cos(\theta_q - \theta_{\mathbf{x}_0}) \cdot \nabla_{\mathbf{x}_1} \{ h_{\sigma} \}(\mathbf{x}_0) + \sin(\theta_q - \theta_{\mathbf{x}_0}) \cdot \nabla_{\mathbf{x}_2} \{ h_{\sigma} \}(\mathbf{x}_0). \tag{3.9}
$$

The collection of HOG operators  $G_{\sigma,q}$  provides a MF representation oriented with  $\theta_{x_0}$  and containing eight redundant frame components (see Section 4.3 of Chapter 2). Whereas the response maps  $h_{\sigma}(x)$  depend linearly on the texture image  $f(x)$ , it is not anymore the case for the HOG operators due to the local alignment of the angle: the HOG operators are not convolution operators themselves, but take advantage of the convolution framework since they are based on the DoG filters.

For obtaining texture measurements  $\eta$ , the responses of HOG operators can be aggregated over regions *M* using component-wise averages (see Fig. 2.15 of Chapter 2). This allows building scale-wise histograms of oriented gradients, where each bin *q* corresponds to the average response of the response map  $h_{\sigma,q}(x)$  of its corresponding operator  $G_{\sigma,q}$  over M. The properties of HOGs are summarized in Table 3.4.

## *2.2.4. Riesz transform*

A more elegant approach to compute directional transitions between pixel values is to compute them in the Fourier domain instead of using pixel differences in Gaussian-

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 $O$ riented edge detectors  $(1<sup>st</sup>-order Gaussian derivative)$  (2  $\mathbf{F} = \mathbf{F} \mathbf{F} \mathbf{F}$  are obtained for each pixel from the image of the  $m \sqrt{m+1}$ gradient and Laplacian filters are kept  $x_1 - 6$  is  $x_2 - 6$  is  $x_3 - 4 - 2$   $x_3 - 3$   $x_4 - 6$  is  $x_1 - 6$  is  $x_2 - 6$  is  $x_3 - 4 - 2$   $x_5 - 3$ *m* Oriented edge detectors Oriented ridge detectors (2nd-order Gaussian derivative) Gaussian LoG  $\theta_2=\frac{\pi}{6}$  $\theta_1 = 0$   $\theta_2 = \frac{\pi}{6}$   $\theta_3 = \frac{\pi}{3}$  $=\frac{\pi}{3}$   $\theta_4 = \frac{\pi}{2}$  $\frac{\pi}{2}$  θ<sub>3</sub> =  $\frac{2\pi}{3}$  θ<sub>3</sub> =  $\frac{5\pi}{6}$  θ<sub>1</sub> = 0 θ<sub>2</sub> =  $\frac{\pi}{6}$  θ<sub>3</sub> =  $\frac{\pi}{3}$  θ<sub>4</sub> =  $\frac{\pi}{2}$  θ<sub>3</sub> =  $\frac{2\pi}{3}$ 3  $\theta_0 = \frac{5\pi}{6}$  $\frac{5\pi}{6}$   $\theta_1 = 0$   $\theta_2 = \frac{\pi}{6}$   $\theta_3 = \frac{\pi}{3}$   $\theta_4 = \frac{\pi}{2}$   $\theta_5 = \frac{2\pi}{3}$   $\theta_6 = \frac{5\pi}{6}$  $\theta_5 = \frac{2\pi}{3}$   $\theta_6 = \frac{5\pi}{6}$   $\theta_1 = 0$   $\theta_2 = \frac{\pi}{6}$   $\theta_3 = \frac{\pi}{3}$   $\theta_4 = \frac{\pi}{2}$   $\theta_5 = \frac{2\pi}{3}$   $\theta_6 = \frac{5\pi}{6}$ 3  $\theta_4=$ 2  $\theta_3 = \frac{\pi}{3}$  $\theta_2=\frac{\pi}{6}$  $\theta_1=0$   $\theta_2=\frac{6}{6}$ *f*(*x*) *x*0

 $g(f(\bm{x}), \bm{m}) = g(f(\bm{R}_{\theta_0}\bm{x}), \bm{m})$ Biome of the strict of perfectors and Aggregation Functions  $\bm{m}$ 

Figure 3.3: Top row: the 38 operators of the MR8 filterbank. At a given position  $x_0$ , the collection of operators  $G_{n=1,...,8}$ { $f$ }( $x_0$ ) consists of a subset of the six directional operators with maximum response over all orientations  $\theta_{1,\dots,6}$  plus the two circularly symmetric operators. This results in eight responses for each position (hence the name of the approach), which are marked in red in the example above.

smoothed images as presented in Eq. (3.8). This also provides the opportunity to easily compute higher-order image derivatives of order *l* as

$$
\frac{\partial^l}{\partial x_d^l} f(\mathbf{x}) \stackrel{\mathcal{F}}{\longleftrightarrow} (j\omega_d)^l \hat{f}(\omega), \tag{3.10}
$$

where  $1 \le d \le D$ . It can be noticed that differentiating an image along the direction  $x_d$  only requires multiplying its Fourier transform by  $j\omega_d$ . Computing  $l^{\text{th}}$ -order derivatives has an intuitive interpretation (*e.g.*, texture gradient for  $l = 1$ , curvature for  $l = 2$ ), which makes them attractive for understanding the meaning of the texture measures in a particular medical or biological applicative context. However, a pure image derivative filter as computed in Eq. (3.10) is high-pass (because multiplied by  $\omega_d$ ) and accentuates high frequencies along  $x_d$ . Therefore, it is desirable to implement image derivatives as all-pass filters, which is provided with the real Riesz transform<sup>8</sup>  $\mathcal{R}\{f\}(\mathbf{x})$ 

8The Riesz transform is the multi-dimensional extension of the Hilbert transform.

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## Table 3.4: HOG properties.



as [64]

$$
\mathcal{R}\{f\}(\boldsymbol{x}) = \begin{pmatrix} \mathcal{R}_1\{f\}(\boldsymbol{x}) \\ \vdots \\ \mathcal{R}_D\{f\}(\boldsymbol{x}) \end{pmatrix} \xrightarrow{\mathcal{F}} - j \frac{\omega}{\|\omega\|} \hat{f}(\omega), \tag{3.11}
$$

It can be noticed by dividing the Fourier representation with the norm of  $\omega$  transforms Eq. (3.10) in *D* all-pass operators  $\mathcal{R}_d$ . For a fixed order *L*, the collection of higher-order all-pass image derivatives are defined in Fourier as

$$
\widehat{\mathcal{R}}^l\{f\}(\omega) = (-j)^L \sqrt{\frac{L!}{l_1! \cdots l_D!}} \frac{\omega_1^{l_1} \cdots \omega_D^{l_D}}{\left(\omega_1^2 + \cdots + \omega_D^2\right)^{L/2}} \widehat{f}(\omega),\tag{3.12}
$$

which yields a total of  $\binom{L+D-1}{D-1}$ *D*−1  $\int_{0}^{1} = \frac{(L+D-1)!}{L!(D-1)!}$  all-pass filters for all combinations of the elements  $l_d$  of the vector *l* as  $|\boldsymbol{l}| = l_1 + \cdots + l_D = L$ . The collection of Riesz operators of order *L* is denoted by  $\mathcal{R}^L$ . A set of band-pass, multi-scale and multi-orientation operator functions  $g_{\sigma,l}(x)$  can be obtained by simply applying the Riesz transform to circularly symmetric wavelets or multi-scale filters, *e.g.*, the LoG filter  $g_{\sigma}$  (see Eq. 3.5)



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Figure 3.4: Two-dimensional  $L^{\text{th}}$ -order Riesz texture operators with functions  $g_{\sigma,l}(x)$ to a  $I$ <sup>th</sup> order image derivatives at a fixed socker  $\frac{1}{\epsilon}$ providing  $L<sup>th</sup>$ -order image derivatives at a fixed scale  $\sigma$ . Applying  $L = 1$  texture operators with functions  $g_{\sigma 10}$  and  $g_{\sigma 01}$  to the input texture f  $\alpha$  edge profiles can be obtained by  $\alpha$ , but much richer profiles can be obtained by linear combinations of  $\alpha$ response maps  $h_{\sigma,1,0}(\mathbf{x})$  and  $h_{\sigma,0,1}(\mathbf{x})$  decomposing f into its v rections, respectively. Qualitatively,  $L = 1$  corresponds to gradient estimation whereas  $\epsilon$  consequent with  $\epsilon$  multimatives,  $\epsilon$  $L = 2$  estimates the Hessian. They both have intuitive in slope for  $L = 1$ , curvature for  $L = 2$ ), which makes them attractive for understanding multiscale Riesz coefficients. An example showing healthy and fibrosis tissues tissues tissues tissues tissues reaning of the texture measures in a particular medical  $\mathbf{v}$  $i$ response maps  $h_{\sigma,1,0}(\boldsymbol{x})$  and  $h_{\sigma,0,1}(\boldsymbol{x})$  decomposing j  $L = 2$  estimates the Hessian. They both have intu  $\frac{1}{2}$  for  $\frac{1}{2}$ ,  $\frac{1}{2}$ ,  $\frac{1}{2}$  can value for  $\frac{1}{2}$ the meaning of the texture measures in a particular medical or biological applicative context.  $C$  $O$  $HCA$ . erators with functions  $g_{\sigma,1,0}$  and  $g_{\sigma,0,1}$  to the input texture *f* with convolution yields response maps  $h_{\sigma,1,0}(\mathbf{x})$  and  $h_{\sigma,0,1}(\mathbf{x})$  decomposing *f* into its vertical and horizontal di- $L = 2$  estimates the Hessian. They both have intuitive interpretations (*e.g.*, texture  $\text{Cohical}$ 

represented in terms of their Riesz components with *N*=2 is depicted in Fig. 2 a). represented in terms of their Riesz components with *N*=2 is depicted in Fig. 2 a). as

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#### $g_{\sigma,l}(x) = \mathcal{R}^l\{g_{\sigma}\}(x).$  $\overline{a}$  $\delta\sigma$ , the Riesz transformation is applied to each  $\delta\sigma$ , the Riesz transformation is applied to each  $\delta\sigma$ .

Examples of 2D real Riesz operators and their application to shown in Fig. 3.4. They are implemented as a plugin for qu  $\frac{1}{2}$  respectively. The Riesz components, which is not desirable for desirable for desirable for desirable for desirable for  $\frac{1}{2}$ on the ePAD radiology platform (see Chapter 13). 3D real Riesz filters are depicted rotation of π*/*2 will switch the responses of *h*<sup>1</sup> and *h*<sup>2</sup> for *N*=1. To ensure that the *z*. *z*, *z* or enapter *z* and are impremented in the γα Examples of *2D* real Riesz operator in Fig. 2.12 of Chapter 2 and are platform<sup>9</sup> (see Chapter 12). block, and every Riesz component *n*= 1*,...,N*+1 is mapped to a multiscale Examples of 2D real Riesz operators and their application to a directional texture are shown in Fig. 3.4. They are implemented as a plugin for quantitative image analysis in Fig. 2.12 of Chapter 2 and are implemented in the QuantImage radiomics web  $\frac{1}{\sqrt{2}}$ 

 $\overline{\mathrm{Diag}}$  operators as defined in  $\overline{\mathrm{E}}$  $r_{\text{NCSZ}}$  operators as defined in Eq. *However, local rotation-invariance/equivariance an* achieved in a convenient fashion through the most interesting property of Riesz tex-Riesz operators as defined in Eq. (3.12) are not locally rotation-invariant/equivariant. However, local rotation-invariance/equivariance and rich MF representations can be ture operators, which is *steerability*. The Riesz operator functions  $g_{\sigma,l}(x)$  are steerable, which will be detailed in 2D in the following text. 2D steerability means that the responses of  $g_{\sigma,l}$  rotated by an angle  $\theta_0$  can be very efficiently computed with a linear

 $9$ https://radiomics.hevs.ch, as of March 1 2017.

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 $g_{\sigma_{i},1,0}$  ( $R_{\theta_0}$ , $n_i$ ,  $0$ ) $R_{\theta_0}$ ,  $\alpha$  ( $\theta$ ),  $\theta$ ,  $\theta$  ( $\theta$ ),  $\theta$ ),  $\theta$ ),  $\theta$ ),  $\theta$ ),  $\theta$  ( $\theta$ ),  $\theta$ ),  $\theta$  ( $\alpha$ ,  $\theta$ ),  $(x,0)$ 



Figure 3.5: Top row: Example of the steering of the first-order Riesz operator  $g_{\sigma,1,0}$ at the position  $x_0 = 0$  with an angle  $\theta_0 = \frac{5\pi}{6}$ . Bottom row: rich MF representations of  $f(x)$  can be obtained from locally steered Riesz filterbanks (exemplified for  $L = 1$ ).

combination parameterized by a steering matrix  $A_{\theta_0}^L$  as

$$
\mathcal{R}^L\{g_{\sigma}\}(\mathbf{R}_{\theta_0}\mathbf{x}) = \mathbf{A}_{\theta_0}^L \mathcal{R}^L\{g_{\sigma}\}(\mathbf{x}),\tag{3.13}
$$

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where  $g_{\sigma}$  is the circularly symmetric function used to control the spatial support of the operators. For order 1 (*i.e.*,  $L = 1$ ),  $A_{\theta_0}^1$  is equal to the 2D rotation matrix  $\mathbf{R}_{\theta_0}$ . For each scale  $\sigma$ , rich *D*-dimensional MF texture representations can be obtained by finding the local angle  $\theta_{x_0}$  maximizing the response of one chosen Riesz operator function  $g_{\sigma,l_1,...,l_D}$ for each position  $x_0$  using steerability [5, 18, 15, 7, 11]. For instance, 2D angle maps corresponding to a texture function  $f$  can be obtained for  $L = 1$  as

$$
\theta_{x_0} := \underset{\theta_0 \in [0, 2\pi)}{\arg \max} \Big( \cos \theta_0 \cdot g_{\sigma, 1, 0} * f(x_0) + \sin \theta_0 \cdot g_{\sigma, 0, 1} * f(x_0) \Big). \tag{3.14}
$$

This is a particular case of the angle alignment framework introduced in Eqs. (3.3) and (3.4).

A collection of MF texture measurements  $\eta$  can be obtained from component-wise averages of the absolute values (or energies) of the Riesz wavelet coefficients steered with angle maps obtained from *e.g.*, Eq. (3.14). Steerability of the real Riesz transform and its use for the construction of MF representations is illustrated in Fig. 3.5. The properties of real Riesz-based texture analysis are summarized in Table 3.5.

By slightly modifying the definition of the Riesz transform (see Eq. 3.12) into its complex form, it is also possible to obtain two-dimensional texture descriptors that can linearly quantify the amount of local circular frequencies. The latter are implicitly modeled by LBP operators, which unfortunately require a binarization and discretiza-

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∀*a* ∈ R*+*, *R f*

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*a* The *n*th-order complex Riesz transform *R<sup>n</sup>* is defined as the *n*-fold iterate of the complex Riesz transform *R*. In the

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Figure 3.6: Impulse responses of Circular Harmonics (CH) of order  $l = 0, \ldots, 5$ . *<sup>s</sup>*,*<sup>k</sup>* for *<sup>n</sup> <sup>=</sup>* 0,..., 5. Top, middle and bottom rows

This is not the case when using the complex Riesz transform. The  $2D$  *l*-th order complex Riesz transform  $\mathcal{R}_\mathbb{C}^l$  is defined in polar coordinates in Fourier as [62] tion of the circular neighborhoods resulting in a potentially large loss of information.

$$
\widehat{\mathcal{R}^l}_{\mathbb{C}}\{f\}(\rho,\vartheta) = e^{jl\vartheta}\widehat{f}(\rho,\vartheta). \tag{3.15}
$$

Using a process similar to construct real Riesz filters, applying the complex Riesz transform to circularly symmetric filters or wavelets (e.g., LoG filters, Eq. 3.5, or circularly symmetric Simoncelli wavelets, Eq. 2.2 of Chapter 2) yields steerable, complex, ability [62], which implies that every steerable representation  $g_{\sigma,L,c}$  with  $c = 1, \ldots, C$  $\frac{1}{\sqrt{2}}$  as and band-pass Circular Harmonic (CH) filters or wavelets  $g_{\sigma,l}(x) = \mathcal{R}_C^l[g_{\sigma}](x) \in \mathbb{C}$ with  $l = 0, \ldots, L$  [32, 38]. CHs stand out as the canonical representation of steercomponents per scale can be obtained with a  $C \times (L + 1)$  complex shaping matrix U as

$$
\begin{pmatrix} g_{\sigma,L,1} \\ \vdots \\ g_{\sigma,L,C} \end{pmatrix} = \mathbf{U} \begin{pmatrix} g_{\sigma,0} \\ \vdots \\ g_{\sigma,L} \end{pmatrix} . \tag{3.16}
$$

As a consequence, any steerable representation can be obtained with a specific shaping matrix U (e.g., gradient and Hessian real Riesz, Simoncelli's steerable pyramid). Examples of CH filters are shown in Fig 3.6. Rich and locally rotation-invariant de-<br>corintions of the LOIDs can be obtained with a year open computational cost by scriptions of the LOIDs can be obtained with a very cheap computational cost by of CH with various orders  $t = 0, 1, \ldots, L$  defines an orthonormal system corresponding to a Fourier basis for circular frequencies up to a maximum order *L*. The process is similar to LBP as presented in [1] but with the desirable properties of a fully linear computing the complex magnitudes of the response maps  $h_{\sigma,l=0,\dots,L}(x)$ . The collection of CH with various orders  $l = 0, 1, \ldots, L$  defines an orthonormal system correspond-

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Table 3.5: Properties of real and complex Riesz texture representations, as well as Steerable Wavelet Machines (SWM).



approach (*e.g.*, no gray-level transformation required) besides the final step computing the complex magnitudes. However, absolute values of CH filters do not define MF representations since the inter-harmonic phase is lost by taking the magnitude of the responses of the operators. Rich MF representations can be obtained by steering all harmonics with a unique local orientation  $\theta_{x_0}$ , as proposed in [16]. Although more computationally expensive than using the magnitudes of the operators, steering them is relatively cheap because CH are self-steerable, resulting in block-diagonal steering matrices  $A_{\theta_0}^L$  for multi-order representations.

Texture measures  $\eta$  can be obtained from the averages or covariances of absolute values (or energies) of steered or non-steered Riesz response maps over a region of interest *M* [7]. The properties of complex Riesz-based texture analysis are summarized in Table 3.5.

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## 2.3. Learned filters

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All texture operators described and discussed in the previous sections are handcrafted. This means that the type of texture information extracted by theses operators is assumed to be relevant for the texture analysis task in hand. Therefore, the design of these operators in terms of the coverage of image scales and directions was based on prior assumptions (*e.g.*, ad-hoc or based on theoretic guidelines, see Section 3.1 of Chapter 1). Whereas classical approaches use machine learning on top of handcrafted representations, more recent approaches proposed to derive the design of the operators from data to identify the combinations of scales and directions that are optimal for the texture analysis task in hand. When compared to handcrafted operators, learned ones reduce the risk of unnecessary modeling of texture properties that are not related to the targeted application (see Fig. 2.5 of Chapter 2). They eliminate human bias to include or exclude arbitrary operator scales and directions. It is worth noting that learned approaches still predominantly use handcrafted components of the aggregation function (*e.g.*, ReLU, sigmoid, pooling, see Section 2.3.3). Three important approaches for learning convolutional texture filters are discussed in this section, including Steerable Wavelet Machines (SWM, Section 2.3.1), Dictionary Learning (DL, Section 2.3.2), and deep Convolutional Neural Networks (CNNs, Section 2.3.3).

#### *2.3.1. Steerable Wavelet Machines (SWM)*

In [12, 17, 14] and [16], we proposed to use SVMs to learn optimally discriminant linear combinations of real or complex Riesz operators. The most interesting property of this approach called *Steerable Wavelet Machines* (SWM) is to combine the flexibility of learned representations with steerability<sup>10</sup>, *i.e.*, we learned the lines  $u_c$  of the shaping matrix  $U$  (see Eq. 3.16) using one-versus-all classification configurations. This yields class-specific steerable signatures of the essential stitches of biomedical tissue (allowing locally rotation-invariant descriptions of the LOIDs) that can be used for building data-specific MF representations. This approach is based on handcrafted representations of image scales based on dyadic circularly symmetric wavelet functions and we are currently extending the learning to image scales as well. The learned operators are limited to the span of Riesz representations. However, it was observed that a limited number of circular harmonics (*e.g.*,  $L = 3, \ldots, 10$ ) yields optimal texture representations for classification. In addition, the span of CH becomes less and less restricted with large values of *L* and can represent any function for  $L \rightarrow \infty$  since it can be interpreted as a Fourier transform for circular frequencies. The properties of SWMs are summarized in Table 3.5.

<sup>10</sup>Linear combinations of steerable subspaces are themselves steerable.

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Figure 3.7: Dictionary learning (DL) process from one single image *fi*(ξ). A collection of 3  $\times$  3 patches  $\gamma_p(\xi)$  are extracted from  $f_i$ . The patch matrices are vectorized to create vectors  $\gamma_p$ . K-means clustering is used in the space spanned by instance vectors  $\gamma_p$  to find *N* atoms  $g_{n,f_i}(\xi)$  that are representative of  $f_i$ . These atoms are called *textons* and can be further used as texture operator functions.

## *2.3.2. Dictionary Learning (DL)*

Recent popular approaches propose to fully learn operators from data (*e.g.*, a collection of bounded discretized texture functions  $I = \{f_1(\xi), f_2(\xi), \ldots, f_I(\xi)\}\)$  with very little constraints besides their maximal spatial support  $G_1 \times \cdots \times G_D$ . A first notable example is unsupervised Dictionary Learning (DL). The learned set of operators will depend on the learning criteria used [61]. Basic dimensionality reduction methods such as Principal Component Analysis (PCA), Independent Component Analysis (ICA), or K-means clustering based on the |*G*| pixel values of  $G_1 \times \cdots \times G_D$  patches  $\gamma_p(\xi)$  can be used to derive *N* essential *atoms*  $g_{n,I}(\xi)$ , which can be further used as operator functions [54]. PCA requires that the atoms are orthogonal to each other (*i.e.*, it removes correlation between them). ICA minimizes the correlation as well as higher order dependence between atoms, which are not necessarily orthogonal. K-means finds *N* prototype atoms  $g_{n,I}(\xi)$  from clustering of the  $|G|$ -dimensional space spanned by the pixel values of the patches  $\gamma_p(\xi)$ . In this particular case, the atoms are also called *textons* [37, 72], which were identified as the elementary units of pre-attentive human texture perception [33]. Textons relate to texture primitives (see Section 2.4 of Chapter 1) and to the LOIDs (see Section 4.1 of Chapter 2). The process for extracting  $3 \times 3$  textons from a texture function  $f(\xi)$  is illustrated in Fig. 3.7.

Another successful unsupervised approach was based on creating a collection of

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atoms from which we can reconstruct the patches  $\gamma_p(\xi)$  with a coefficients vector  $\alpha_p$ containing a minimum number of non-zero elements [41, 24]. Every atoms  $g_{n,I,\lambda}$  are vectorized, transposed and piled up to create the  $|G| \times N$  dictionary matrix **D** with N atoms. Solving the following optimization problem for a collection of *P* patches can be used to compute both **D** and  $\alpha$  as

$$
\mathbf{D} := \underset{\mathbf{D}, \alpha}{\arg \min} \sum_{p=1}^{P} ||\mathbf{\gamma}_p - \mathbf{D}\alpha_p||_2^2 + \lambda ||\alpha_p||_1,
$$
 (3.17)

where  $\|\cdot\|_1$  is the  $\ell_1$ -norm<sup>11</sup> and  $\lambda$  is controlling the sparsity of the atom coefficients  $\alpha$ . Additional topological constraints were added in Topographic Independant Component Analysis (TICA) to take into account the important knowledge that patches that are spatially close to each other have similar statistical propoerties [31]. All texture operators dictionaries D learned with unsupervised approaches (*e.g.*, K-means, PCA, ICA, TICA, reconstruction, see Eq. 3.17) are not directly optimizing a discriminative criteria, which means that they are not necessarily optimal for biomedical texture classification of a set of considered tissue types. To tackle this issue, supervised dictionary learning was proposed [41], where D is obtained as

$$
\mathbf{D} := \underset{\mathbf{D},b,\alpha}{\arg\min} \sum_{p=1}^{P} C(y_p, \gamma_p^T \mathbf{D} \alpha_p + b) + \lambda ||\mathbf{D}||_F^2, \tag{3.18}
$$

where  $\gamma_p$  are the vectorized training patches and their labels  $(y_p \in \{-1, +1\})_{p=1,\dots,P}$ ,  $C(y_p, \breve{y}_p)$  is the cost function<sup>12</sup>, *b* the bias, and  $|| \cdot ||_F$  the Frobenius matrix norm<sup>13</sup>. An inherent challenge of DL methods is the large number of free parameters (*i.e.*, features or variables) to learn which is equal the size (*i.e.*, measure) of the dictionary  $|G| \times N$ , resulting in a very high-dimensional feature space. This typically requires a very large collection of training patches to respect the recommended ratio between feature dimensionality and number of training instances equal to ten [28]. As an order of magnitude, 121,0000 training patches are required to learn 1000  $11 \times 11$  atoms. A collection of N texture measurements  $\eta$  can be obtained by computing the averages of the absolute values or energies of the response maps  $h_n(\xi) = (g_n * f)(\xi)$  within a region of interest *M*. The properties of DL-based texture analysis are summarized in Table 3.6.

<sup>&</sup>lt;sup>11</sup>The  $\ell_1$ -norm of a vector *x* is  $||x||_1 = \sum_{d=1}^D |x_d|$ , whereas the  $\ell_2$ -norm is  $||x||_2 = \sqrt{\sum_{d=1}^D x_d^2}$ . <sup>12</sup>The cost function defines how label prediction errors of the estimations  $\check{y}_p$  are penalized. The logistic

loss function is used in [41] and is defined as  $C(y_p, \tilde{y}_p) = \log(1 + e^{-y_p \tilde{y}_p})$ .

<sup>&</sup>lt;sup>13</sup>The Frobenius matrix norm is defined as  $||A||_F = \sqrt{\sum_{i=1}^{I} \sum_{j=1}^{J} a_{i,j}^2}$ , where  $a_{i,j}$  are the elements of **A**.

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#### *2.3.3. Deep Convolutional Neural Networks (CNN)*

One major and very successful work on filter learning are CNNs and their deep architectures<sup>14</sup> [26, 67, 35, 55, 56]. Review and applications of deep learning in texture analysis for tissue image classification are further detailed in Chapters 4, 9 and 10. In a nutshell, deep CNNs consist of a cascade of *Q* convolutional layers, where each of the latter typically contains

- (i) a multi-channel convolution of the input  $f_{n',q}(\xi)^{15}$  with a set of  $N G_1 \times \cdots \times G_D$ multi-channel operator functions *gn*,*n*′,*q*(ξ), called *receptive fields*,
- (ii) a simple pointwise non-linear gating  $V(x)$  of the response map  $h_{n,q}(\xi)$ ,
- (iii) an optional cross-channel normalization [35] and,
- (iv) a pooling operation resulting in a down- or up-sampling of  $h_{n,q}(\xi)$ .

<sup>14</sup>A notable exception is the Scattering Transform (ST) [3], which is a hancrafted deep CNN. <sup>15</sup>The first layer  $q = 1$  will use the original image  $f_{n',1}(\xi)$  as input, whereas the next layers  $q = 2, \ldots, Q$  will use the  $n' = 0, \ldots, N'$  outputs (*i.e.*, *channels*)  $f_{n',q-1}(\xi)$  of the previous layer as input for the convolution.

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In 2D, the multi-channel convolution in a layer *q* between the input  $f_{n',q-1}(\xi)$  and the set of *N* multi-channel filters  $g_{n,n',q}(\xi)$  is

$$
h_{n,q}(\xi) = \sum_{n'=1}^{N'} (g_{n,n',q} * f_{n',q-1})(\xi).
$$
 (3.19)

(3.19) yields a multi-channel response map  $h_{n,q}(\xi)$  with a dimensionality of  $F_1 \times F_2 \times N'$ for a 2D input function of domain *F*. The convolution is two-dimensional, where no cross-channel convolution is carried out. Popular examples of non-linear gating functions  $V(x)$  are the Rectified Linear Unit (ReLU)

$$
\mathcal{V}_{\text{ReLU}}(x) = \max(0, x),
$$

the sigmoid function

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$$
\mathcal{V}_{\text{sig}}(x) = \frac{1}{1 + e^{-x}},
$$

the absolute value and the energy. The aim of the pooling operation is to down- or up-sample feature maps to achieve a multi-scale analysis throughout the cascade of convolutional layers. Whereas the upsampling operation is often straightforward (no specific super-resolution approach [55]), the downsampling requires using a criteria on which value to keep over local patches (*e.g.*, maximum, minimum, median, sum, average). The structure of one convolutional layer is depicted in Fig. 3.8.

Deep CNN architectures consist of a cascade of convolutional layers containing a large collection of operator functions  $g_{n,n',q}(\xi)$  learned to minimize a cost function  $C(y_p, \breve{y}_p)$ . The total number of free parameters is equal to the product of the number of channels per operator N', the number of operator per layer N, the size of the spatial support  $|G|$  of the operators, and the number of layers Q, if we assume that N, N' and |*G*| are identical over all layers. It is very common for CNN architectures designed for classification to have a final *Fully Connected* (FC) layer that computes the decision value based on a linear combination of all elements of the final feature maps  $h_{n,Q}(\xi)$ . As an example, the complete forward function  $u_{forward}(f(\xi))$  of a classification CNN transforms the input texture  $f_p(\xi)$  into an estimated class label  $\tilde{y}_p^{16}$  is a composition of all functions  $u_q$  from all layers as

$$
u_{\text{forward}}(f(\xi); \mathbf{g}): \quad \mathbf{F} \to \mathbb{R}, \tag{3.20}
$$

$$
\breve{y}_p = u_{\text{forward}}(f_p(\xi); \mathbf{g}) = u_{\text{FC}}(\cdots u_2(u_1(f_p(\xi); \mathbf{g}_1); \mathbf{g}_2) \cdots); \mathbf{g}_{\text{FC}}), \tag{3.21}
$$

where  $g_q$  is the collection of free parameters of the operator functions  $g_{n,n',q}(\xi)$  in the layer *q*, and *g* is the total collection of free parameters of the model. When CNN ar-

<sup>&</sup>lt;sup>16</sup> $\tilde{y}_p$  can be binary (*e.g.*,  $\tilde{y}_p \in \{-1, +1\}$ ) or continuous (*e.g.*, probability  $\tilde{y}_p \in [0, 1]$ , or  $\tilde{y}_p \in \mathbb{R}$ ). We will assume that  $\breve{y}_p \in \mathbb{R}$  in our example.

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Figure 3.8: Structure of one typical convolutional layer of CNNs. The multi-channel output of the previous layer  $q - 1$  constitutes the input  $f_{n',q-1}(\xi)$  of the layer q. A multichannel convolution is carried out between the channels of *fn*′,*q*−<sup>1</sup> and the collection of  $n = 1, \ldots, N$  operator functions  $g_{n,n',q}$  and yields a collection of response maps  $h_{n,q}(\xi)$ as detailed in Eq. (3.19). The latter undergo a simple pointwise non-linear gating  $V(x)$ , followed by a pooling operation to output the final feature maps  $f_{n,q}(\xi)$ . In the example above, the 5  $\times$  5 operator functions  $g_{n,n',q}$  are initialized with random values following a normal distribution with zero mean. The non-linear gating function  $\mathcal V$  is a ReLU, and the pooling operation is a  $4\times$  downsampling process (noted as  $\downarrow$  4) where the maximum value is kept over  $4 \times 4$  patches and a distance (or *stride*) of 4 between their centers.

chitectures are designed for segmentation (*e.g.*, U-Net [55]), the output of the forward function is an estimated segmentation map  $\tilde{y}_p(\xi)$ . Based on a set of training textures from which the true labels  $y_p$  are known, it is possible to compute the prediction errors between  $\breve{y}_p$  and  $y_p$ . The cost of this error is determined by the cost function  $C(y_p, \breve{y}_p)$ . Typical  $C(y_p, \breve{y}_p)$  (also called *loss* function) are the logistic loss function for classification, or Euclidean loss for regression. The total cost function  $C_{\text{tot}}$  over the entire

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Figure 3.9: CNN architecture with a cascade of  $Q = 5$  convolutional and one fully connected layers. The forward function  $u_{forward}(f(\xi); g)$  is a composition of layer-wise functions  $u_g$  and transforms an input image  $f_p(\xi)$  into a predicted label  $\check{y}_p$ . The cost function  $C_{\text{tot}}$  of accumulated errors over the entire training set is minimized through several epochs using backpropagation to update the free parameters *g*. The latter allows learning the optimal profile of the convolutional operator functions  $g_{n,n',q}$ .

training set (called *epoch*) is accumulating errors over instances *p* as

$$
C_{\text{tot}}(\mathbf{g}) = \frac{1}{P} \sum_{p=1}^{P} C\left(y_p, u_{\text{forward}}(f_p(\mathbf{\boldsymbol{\xi}}); \mathbf{g})\right).
$$

After computing the total cost over one epoch  $t$ , the parameters of the model  $g_t$  are modified to minimize the total loss as

$$
\boldsymbol{g}_{t+1} = \boldsymbol{g}_t - \beta_t \frac{\partial C_{\text{tot}}}{\partial \boldsymbol{g}} (\boldsymbol{g}_t),
$$
 (3.22)

where  $\beta_t$  is the learning rate at the epoch *t*. Solving Eq. (3.22) learns the operator functions  $g_{n,n',q}(\xi)$  and is called *backpropagation*.  $\frac{\partial C_{\text{tot}}}{\partial g}$  is computed using the chain rule, which allows combining the derivatives of the loss with respect to the parameters  $g_q$  of each consecutive layer [26]. Solving Eq. (3.22) through a total number *T* of epoch can be efficiently carried out with gradient descent optimization techniques. An example of a CNN architecture, its forward function and backpropagation is illustrated in Fig. 3.9. CNN-based texture analysis do not yield texture measurements *per se* as it directly outputs a tissue class probability or segmentation map. The aggregation function results from a cascade of pooling operations and, if applicable, the FC

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## Table 3.7: Properties of deep CNNs for biomedical texture analysis.



layer. The number of free parameters can be extremely large. For instance, a network with 10 layers, 10 11  $\times$  11 operators per layer with 10 channels and with a 50  $\times$  50 FC layer leads to  $> 10^8$  free parameters. It therefore requires very large training sets to correctly learn the model, which is most often difficult to get in biomedical imaging. An approach called *transfer learning* is often used is to tackle this problem [53, 46] (see Section 4.3.3 of Chapter 4). It consists of re-using models trained with millions of images from other domains (*e.g.*, photographs in ImageNet [10]). In the particular case of biomedical texture analysis, reusing these models is risky because the types of invariances learned by networks based on photographic imagery resulting from scene captures obtained with varying viewpoints is very different from the ones desirable in BTA (see Section 3.3 of Chapter 1). In interesting observation is that when large datasets are available, the type of operators learned in the first layers of deep CNNs share very similar properties with handcrafted convolutional filters such as LoG, Gabor, MR8, and Riesz. The properties of CNNs for biomedical texture analysis are summarized in Table 3.7.

## *2.3.4. Data augmentation*

Data augmentation aims to tackle two important limitations of filter learning approaches with a large number of free parameters: DL and CNNs [35, 53, 46, 55]. First, both

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approach require very large training sets to respect an acceptable ratio between the number of free parameters and training instances and allow suitable generalization performance (*i.e.*, limiting the overfitting of the training data). Second, neither of the methods are invariant to local scalings or local rotations. Whereas invariance to image scale is most often not desirable for biomedical texture analysis, invariance to local rotations is fundamental (see Section 3.3 of Chapter 1). Data augmentation consists of generating additional training instances from geometric transformations (including non-rigid) of the available ones, and to further train the models with these new instances. It provided very important performance gain in various applications [35, 53], including biomedical [55]. However, it is obvious that forcing convolutional operators to recognize sheared, scaled and rotated versions of the texture classes without augmenting the degrees of freedom will result in a strong decrease of the specificity of the model. In particular, forcing invariance to local rotations will make operators insensitive to directions, which was illustrated in [25]. A quantitative comparison between circularly symmetric and locally aligned operators (*e.g.*, MF representations) is detailed in Section 4.4 of Chapter 2. In this context, learning steerable filters allows adapting the texture representation while keeping local rotation-invrariance and high specificity of the operators [12, 16].

## 3. Gray-level matrices

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There are mainly three approaches based on Gray-Level Matrices (GLM): Gray-Level Co-occurrence Matrices (GLCM), Gray-Level Run-Length Matrices (GLRLM), and Gray-Level Size Zone Matrices (GLSZM). Their simplicity is at the origin of their popularity and many implementations can be found (*e.g.*, MaZda<sup>17</sup>, see Chapter 11, LIFEx<sup>18</sup>, Matlab<sup>19</sup>, Scikit-image<sup>20</sup>, QuantImage web platform<sup>21</sup>, see Chapter 12, ePAD texture plugins<sup>22</sup>, see Chapter 13). They are often used together. They are based on discrete texture functions  $f(\xi)$  and operators  $G_n$ . Their extensions to 3D are straightforward. However, they suffer from several imperfections. A major one is the nonsystematic coverage and poor preservation of image scales and directions, especially when the spatial supports  $G_{1,n} \times \cdots \times G_{D,n}$  of their operators are large. Therefore, they are best suited for applications where the size of the ROIs are small (*e.g.*, small lesions in low-resolution medical images). They also require drastic reductions of gray levels, where the values of  $f(\xi) \in \mathbb{R}$  are quantized with  $\Lambda$  values as  $f_{\Lambda}(\xi) \in \{1, \ldots, \Lambda\}$ . The

<sup>17</sup>http://www.eletel.p.lodz.pl/programy/mazda/, as of March 4 2017.

<sup>18</sup>http://www.lifexsoft.org, as of November 20 2016.

 $19$ http://www.mathworks.com/help/images/ref/graycomatrix.html, as of November 20 2016.

 $^{20}$ http://scikit-image.org/docs/stable/api/skimage.feature.html, as of February 26 2017. <sup>21</sup>https://radiomics.hevs.ch, as of March 4 2017.

<sup>22</sup>http://epad.hevs.ch, as of March 4 2017.

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quantization is typically based on  $\Lambda = 8, 16, 32$ , which results in an important potential loss of information when analyzing rich image contents coded with 12 to 16 bits<sup>23</sup>. Their properties are reviewed and detailed in the three following subsections 3.1, 3.2 and 3.3.

## 3.1. Gray-Level Co-occurrence Matrices (GLCM)

GLCMs [30] can be seen as a collection of operators mapping the discretized and quantized image function  $f_\Lambda(\xi)$  to binary output values at a position  $\xi_0$  as

$$
\mathcal{G}_{\Delta k}^{\lambda_i, \lambda_j}\{f_{\Lambda}\}(\xi_0) = \begin{cases} 1 & \text{if } f_{\Lambda}(\xi_0) = \lambda_i \text{ and } f_{\Lambda}(\xi_0 + \Delta k \circ \Delta \xi) = \lambda_j, \\ 0 & \text{otherwise,} \end{cases} \tag{3.23}
$$

where  $\lambda_i$  and  $\lambda_j$  are the pixel values at positions  $\xi_0$  and ( $\xi_0 + \Delta k \circ \Delta \xi$ ), respectively. ∆*k* and ∆ξ contains the dimension-wise shifts and sampling steps, respectively (see Section 2.2 of Chapter 1).  $\Delta k \circ \Delta \xi$  denotes the element-wise product<sup>24</sup> between the two vectors as

$$
\Delta \boldsymbol{k} \circ \Delta \boldsymbol{\xi} = \begin{pmatrix} \Delta k_1 \cdot \Delta \xi_1 \\ \vdots \\ \Delta k_D \cdot \Delta \xi_D \end{pmatrix}.
$$

The aggregation function is integrative. It counts the responses of the operators and organizes them in square *co-occurrence* matrices of dimensions  $\Lambda^2$  indexed by  $(\lambda_i, \lambda_j)$ , where  $\Lambda$  is the number of gray-levels. A series of scalar texture measurements<sup>25</sup>  $\eta$ is obtained by computing statistics (*e.g.*, contrast, correlation, entropy) from the cooccurrence matrices. Their properties are summarized in Table 3.8. GLCMs are not invariant to local rotations, but the later is often approximated by either regrouping the counts of operators over all directions in a shared matrix, or by averaging scalar texture measurements from co-occurrences matrices obtained with different directions (see Fig. 3.10).

## 3.2. Gray-Level Run-Length Matrices (GLRLM)

GLRLMs [22] count the number of aligned pixels (called stride) with equal gray-level value λ, length γ ∈ N<sup>\*</sup> and direction θ. Their operators  $G_{\lambda, \gamma, \theta}(f_\Lambda)(\xi_0)$  yield a value of 1 if a stride starting at  $\xi_0$ , of length  $\gamma$ , and direction  $\theta$  is detected, and 0 otherwise. An example of a GLRLM operator and its response map is depicted in Fig. 3.11.

The aggregation function counts the number of strides detected with the corre-

 $^{23}$ Image pixels encoded with 16 bits can take more than 65,000 possible values.

<sup>24</sup>It is also called the *Hadamard* product.

<sup>25</sup>They are often called *Haralick features* [30].

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## Table 3.8: GLCM properties.





Figure 3.10: Approximate local rotation-invariance with GLCMs: the responses of GLCM operators  $G_{\Delta k_1,\Delta k_2}^{\lambda_i,\lambda_j}$  are combined over four directions  $\theta_{1,\dots,4}$  to reduce the directional sensitivity of the operators and approximate locally rotation-invariant texture analysis. In 2D, the angle between the vectors  $\Delta \mathbf{k}$  and  $\mathbf{e}_1 = (1, 0)$  is noted  $\theta$  (see Section 2 of Chapter 2).

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## Table 3.9: GLRLM properties.



sponding operator and organizes them in a *run-length* matrix indexed by  $\lambda$  and  $\gamma$ . The size of this matrix is the number  $\Lambda$  of gray-levels considered times the number of stride lengths tested. The texture measures  $\eta$  are obtained from statistics of the runlength matrices (*e.g.*, short run emphasis, run-length non-uniformity, see [71]). Their properties are summarized in Table 3.9.

## 3.3. Gray-Level Size Zone Matrices (GLSZM)

GLSZMs [60] are extending the concept of GLRLM to zone areas or volumes. Their operators  $G_{\lambda,\zeta} \{f_{\Lambda}\}\xi_0$ ) yield a binary value of 1 if  $\xi_0$  belongs to a uniform zone with gray-level  $\lambda$ , *i.e.*, any zone with area equal to  $\zeta$  containing  $\xi_0$ , and area  $\zeta \in \mathbb{N}^*$ , and  $\xi_0$ does not belong to a uniform zone with larger area.

The aggregation function counts the number of zones detected with the corresponding operator and organizes them in a *size-zone* matrix indexed by  $\lambda$  and  $\zeta$ . The size of this matrix is the number  $\Lambda$  of gray-levels considered times the number of zone ar-

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Figure 3.11: Application of a GLRLM operator  $G_{\lambda=7,\gamma=4,\theta=0}$  to an input image  $f_{\Lambda}$ . Its response map  $h_{\lambda=7,\gamma=4,\theta=0}(\xi)$  highlights the presence of the sought stride with graylevel 7, length 4 and orientation 0 at the position  $\xi_0$ .

Table 3.10: GLSZM properties.

<b>Operator linearity</b>	Non-linear.
<b>Handcrafted</b>	Yes.
3D extension	Straightforward: the search for contiguous pixels with identical gray-level values can be extended from 8-connected 2D neighborhoods to their 26-connected 3D counterparts.
<b>Coverage of image</b> directions	Complete.
<b>Directionality and local</b> rotation-invariance	Insensitive to image directions. Elongated and circular zones are mixed.
<b>Characterization of the</b> <b>LOIDs</b>	No.
<b>Coverage of image</b> scales	Complete if the maximum zone surface $\zeta$ is equal to the area of the image. However, the notion of scale is ill-defined because a fixed zone area $\zeta$ can correspond to both elongated or circular regions.
<b>Band-pass</b>	Qualitatively equivalent in the sense that the mean value of the image is not influ- encing the output value of the operator. It is worth noting that the transfer function is not defined in the Fourier domain because the operator is nonlinear.
<b>Gray-level reduction</b>	A reduction is required to avoid obtaining very large and sparse size-zone matrices. Typical gray-level reductions are $\Lambda = 8, 16, 32$ .
<b>Illumination-invariance</b>	No, although it is approximated by reducing the number of gray-levels $\Lambda$ .
<b>Aggregation function</b>	Integrative: counts the binary responses of each GLSZM operator $G_{\lambda,\zeta}$ over M and organizes them in a corresponding size-zone matrix indexed by integer values of $\lambda$ and $\zeta$ . Scalar texture measurements $\eta$ are obtained from statistics of the size-zone matrices.

eas tested. The texture measures  $\eta$  are obtained from statistics of the matrices, which are the same as for the GLRLM (see [71]) plus two additional measures introduced in [60]. Their properties are summarized in Table 3.10.

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## 4. Local Binary Patterns (LBP)

Rotation-invariant Local Binary Patterns (LBP) were first introduced by Ojala *et al.* in 2002 [45] and many extensions were proposed later on (*e.g.*, [1, 27, 39]). At a position  $\xi_0$  of the image f, the LBP operator  $G_{\gamma,r}$   $(f)(\xi_0)$  describes the organization of binarized pixels over circular neighborhoods  $\Upsilon(\gamma, r, \xi_0)$  of radius *r* containing  $\gamma$  equally spaced points. The gray value of points that do not fall exactly in the center of pixels are estimated by interpolation (see Fig. 3.12, top). The decimal value of the operator is given by the binary sequence of the circular neighborhood (*e.g.*,  $10101010 = 170$ , see Fig. 3.12, middle). The binary value of a point  $p \in \Upsilon(\gamma, r, \xi_0)$  is 1 if  $f(\xi_1)$  $f(\xi_0)$ , and 0 otherwise, with  $\xi_1$  the pixel location where p is located. An example of approximately locally rotation-invariant LBP texture analysis is depicted in Fig. 3.12 middle and bottom.

The responses of the operators are aggregated over a ROI *M* by counting the binary sequences (or decimal values) and organizing them in a histogram. The latter can be used for extracting texture measures  $\eta$ . The properties of LBPs are summarized in Table 3.11.

## 5. Fractals

Another popular method in BTA is to estimate the fractal properties of biomedical tissue, *e.g.*, how structures are similar across a series of monotonously increasing scales [42, 2]. The latter is measured with the *Fractal Dimension* (FD)  $\eta_{FD}$ . The larger the FD, the most regular structures are through multiple scales and corresponds closely to our intuitive notion of roughness [48]. The maximum FD is equal to the dimensionality *D* of the texture function. For 2D texture functions indexed by the spatial coordinates  $(x_1, x_2) \in \mathbb{R}^2$ , we typically consider the set  $S = \{(x_1, x_2) \in \mathbb{R}^2 : f(x_1, x_2) =$ *y*}  $\subset \mathbb{R}^2$  for a fixed value *y*  $\in \mathbb{R}$ . Note that *S* is a level set of *f*. Then, the fractal dimension can take values  $0 \le \eta_{FD} \le 2$ . A comprehensive description and review of fractal analysis for BTA is presented in Chapter 5.

A popular method for computing the FD is the *box-counting* approach, while several other methods exist (*e.g.*, box counting and probabilities [34], based on Fourier analysis [48] or wavelets [49], see Chapter 5 for detailed descriptions and more approaches). Box-counting relies on a collection of multi-scale operators  $G_n$ . For simplification, let us assume that the latter have square or cubic spatial support  $G_{1,n} = G_{2,n}$  $\cdots = G_{D,n} = s_n$ . The collection of operators have monotonously increasing spatial supports  $G_n \subset G_{n+1}$ . In order to simply exemplify the box-counting algorithm, let us consider binary images  $f_\tau(\xi) \in \{0, 1\}$  obtained from the binarization of an input image  $f(\xi)$  with a threshold  $\tau^{26}$ . At a position  $\xi_0$ , each fractal operator  $G_n$  counts the number

 $^{26}$ The binarization step or grey-level reduction is usually not recommended for box-counting as this would

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## Table 3.11: LBP properties.



*p* of pixels with value equals to 1 (*i.e.*,  $f_\tau(\xi_0) = 1$ ) over its support  $G_n$ . This yields *N* response maps  $h_n(\xi_0) = G_n(f_\tau)(\xi_0)$ .

The aggregation function consists of fitting a log-log profile of the averages of *hn* through the responses of the collection of operators with spatial supports of varying sizes  $s_n$ , where  $|G_n| = s_n^D$ . The average number p of pixels with value equals to 1 over a region of interest *M* is computed as

$$
p(s_n) = \frac{1}{|M|} \sum_{\xi \in M} h_n(\xi).
$$

both arbitrarily discard important texture information and degrade the stability of the calculation of  $\eta_{FD}$ (see [2, 57] and Chapter 5).

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Figure 3.12: Extraction of LBP texture measures on  $f_1(\xi)$ . Top row: example of two different circular neighborhoods  $\Upsilon(\gamma, r, \xi_0)$ . Middle row: a LBP operator  $G_{8,1}\{f_1\}(\xi_0)$ encodes the LOIDs at the position  $\xi_0$  over a circular neighborhood  $\Upsilon(8, 1, \xi_0)$  with  $\gamma =$ 8 equally-spaced points and a radius  $r = 1$  pixel. Its binary response for a  $+$ -shaped primitive is 10101010, which correspond to a decimal value of 170. The response maps of the operators are aggregated over the region *M* by counting the binary codes and organizing them into an histogram of texture measurements  $\eta_M$ . Bottom row: local rotation-invariance with LBPs. Local image rotations  $\mathbf{R}_{\theta_0,\xi_0} \cdot \boldsymbol{\xi}$  correspond to bit-wise circular shifts of the binary codes. Local rotation-invariance can be achieved by minimizing the decimal values of the binary codes over all possible discrete circular shifts  $\theta_0 = \frac{q\pi}{4}$ ,  $\forall q \in \{0, 1, ..., 7\}.$ 

The FD is obtained with the value of  $\eta_{FD}$  that best fits the following relation

$$
\log\left(p(s_n)\right) \simeq \eta_{\text{FD}} \log\left(s_n\right). \tag{3.24}
$$

In this particular case,  $\eta_{FD} = 2$  corresponds to a perfectly regular image filled with ones. The box counting method is illustrated in Fig. 3.13, where two biomedical textures with different FD are compared. The properties of fractal-based texture analysis are summarized in Table 3.12 and are further developed in Chapter 5. The FD is invariant to image scale, which entails the risk of regrouping tissue structures of different natures (see Section 3.3 of Chapter 1).

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Figure 3.13: 2D box-counting method used to estimate the fractal dimension  $\eta_{FD}$  of a texture. Although not recommended in general, a binarization of the texture images is used in this example. The latter is carried out with a thresholding operation,where the threshold  $\tau$  was chosen to highlight cell nucleus in  $f_1$  and the vascular structure in  $f_2$ . The visualization of the profiles  $p(s_n)$  reveals the fractal nature of the vascular structure with a FD of 1.8. The maximum FD bound of 2 corresponding to a fully white image corresponds to the dashed red line in the profiles  $p(s_n)$ .

## 6. Discussions and conclusions

A qualitative comparison of most popular BTA approaches was proposed under the light of the general framework introduced in Section 3.1 of Chapter 1, and based on the comparison dimensions presented in Chapter 2. Our aim is to both provide a user guide for choosing a BTA method that is relevant to the problem in hand, as well as to provide insights on key aspects required to build the next generation of BTA approaches. The review focused on most popular group of methods and was not exhaustive. It included (i) convolutional methods and their operator subtypes: circu-

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Table 3.12: Properties of fractal-based texture analysis using the box-counting method.



larly/spherically symmetric, directional and learned, (ii) gray-level matrices and their subtypes: GLCMs, GLRLMs, GLSZMs, (iii) local binary patterns and a few extensions, and (iv) fractals based on the box-counting method. All approaches' operators were found to be equivariant to translations, which is an inherited property from the general framework introduced in Section 3.1 of Chapter 1. However, very few methods were combining both the ability to characterize the LOIDs and with invariance to local rotations. In particular, most approaches are either invariant to local rotations because they are insensitive to directions (*e.g.*, combined GLCM or GLRLM operators, LoGs, fractals) and they cannot characterize the LOIDs or, they are directional but not locally rotation-invariant (*e.g.*, unidirectional GLCM or GLRLM operators, Gabor wavelets, unaligned real Riesz, DL, CNNs). Within this dilemma, one may favor directional versus directionally insensitive approaches depending on the expected importance of the LOIDs versus invariance to local rotations, respectively. The only approaches that are able to characterize the LOIDs with invariance to (local) rigid transformations are LBPs as well as CH and steered real Riesz wavelets. Other factors must be taken into account to evaluate the relevance of the method. A important property which is responsible for the success of approaches such as DL and CNNs is the ability to

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derive the texture operators<sup>27</sup> from the data in a supervised or unsupervised fashion. A challenge for the success of the learning-based methods is to have sufficient representation of each intra-class variants, which are most often not available in focused and innovative biomedical applications (see Chapter 6). Quick fixes have been extensively used to tackle this challenge. A first one is to introduce implicit handcrafted invariances with data augmentation, which has the undesirable effects of both increasing the training computational load (minor issue), and more importantly to decrease the specificity of the model [25] (see Section 2.3.4). A second fix is to use transfer learning in order to recycle models trained on other image types. This is not exempt of risks as borrowing models from general computer vision with strong invariance to scale  $(e.g.,$  models trained on ImageNet) are not fulfilling the requirement of BTA (see Section 2.3). An inherent risk of methods requiring important gray-level reductions or binarization (*e.g.*, GLM, LBP, fractals based on box-counting) is to miss or mix important texture properties. It was found that methods based on learned steerable wavelets (*i.e.*, SWMs [16, 12]) were regrouping several desirable properties such as the ability to learn optimally discriminant sets of LOIDs with invariance to rigid transformation, with a small amount of training data, and without requiring neither data augmentation nor gray-level reductions. Few methods were found to provide easily interpretable texture measurements. Notable examples among them are the contrast and energy measures of GLCMs, the fractal dimension and roughness [48] (see Chapter 5), as well as 2D and 3D steered real Riesz wavelets that are measuring image derivatives of various orders in a locally rotation-invariant fashion [15, 11, 18, 5] (see Section 2.4.4 of Chapter 12).

We recognize several limitations of the current review, including the absence of quantitative comparison of performance, computational complexity and time of the approaches. Detailed reviews of specific BTA properties, approaches or applications are presented in Chapter 7 (invariance to rigid transformations), Chapters 4 and 9(deep learning), Chapter 6 (machine learning), and Chapter 10 (digital histopathology).

## Acknowledgments

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#### Bibliography

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1. Timo Ahonen, Ji? Matas, Chu He, and Matti Pietikinen. Rotation Invariant Image Description with Local Binary Pattern Histogram Fourier Features. In Arnt-Brre Salberg, JonYngve Hardeberg, and Robert Jenssen, editors, *Image Analysis*, volume 5575 of *Lecture Notes in Computer Science*, pages 61–70. Springer Berlin Heidelberg, 2009.

 $27$ CNNs with FC layers are also learning the aggregation function.

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#### 98 BIBLIOGRAPHY

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- 2. Omar. S Al-Kadi and D Watson. Texture analysis of aggressive and nonaggressive lung tumor CE CT images. *IEEE Transactions on Biomedical Engineering*, 55(7):1822–1830, July 2008.
- 3. Joan Bruna and Stephane G. Mallat. Invariant Scattering Convolution Networks . *IEEE Transactions on Pattern Analysis and Machine Intelligence*, 35(8):1872–1886, 2013.
- 4. Andreas Burner, Rene Donner, Marius Mayerhoefer, Markus Holzer, Franz Kainberger, and Georg Langs. Texture bags: Anomaly retrieval in medical images based on local 3D-texture similarity. In Hayit Greenspan, Henning Müller, and Tanveer Syeda-Mahmood, editors, *Medical Content-based Retrieval for Clinical Decision Support*, volume 7075 of *MCBR-CDS 2011*, pages 116–127. Lecture Notes in Computer Sciences (LNCS), September 2012.
- 5. Nicolas Chenouard and Michael Unser. {3D} Steerable Wavelets in Practice. *IEEE Transactions on Image Processing*, 21(11):4522–4533, 2012.
- 6. Özgün Çiçek, Ahmed Abdulkadir, Soeren S Lienkamp, Thomas Brox, and Olaf Ronneberger. 3D *U-Net: Learning Dense Volumetric Segmentation from Sparse Annotation*, pages 424–432. Springer International Publishing, Cham, 2016.
- 7. Pol Cirujeda, Yashin Dicente Cid, Henning Muller, Daniel Rubin, Todd A Aguilera, Billy W Loo ¨ Jr., Maximilian Diehn, Xavier Binefa, and Adrien Depeursinge. A 3–D Riesz–Covariance Texture Model for Prediction of Nodule Recurrence in Lung CT. *IEEE Transactions on Medical Imaging*, 2016.
- 8. Ingrid Daubechies. *Ten lectures on wavelets*, volume 61. SIAM, 1992.
- 9. John G Daugman. Uncertainty relation for resolution in space, spatial frequency, and orientation optimized by two-dimensional visual cortical filters. *Journal of the Optical Society of America. A, Optics and image science*, 2(7):1160–1169, 1985.
- 10. Jia Deng, Wei Dong, R Socher, Li-Jia Li, Kai Li, and Li Fei-Fei. ImageNet: A large-scale hierarchical image database. In *IEEE Conference on Computer Vision and Pattern Recognition*, CVPR 2009, pages 248–255, 2009.
- 11. Adrien Depeursinge, Antonio Foncubierta-Rodríguez, Dimitri Van De Ville, and Henning Müller. Lung texture classification using locally-oriented Riesz components. In Gabor Fichtinger, Anne Martel, and Terry Peters, editors, *Medical Image Computing and Computer Assisted Intervention – MICCAI 2011*, volume 6893 of *Lecture Notes in Computer Science*, pages 231–238. Springer Berlin / Heidelberg, sep 2011.
- 12. Adrien Depeursinge, Antonio Foncubierta-Rodríguez, Dimitri Van De Ville, and Henning Müller. Rotation–covariant texture learning using steerable Riesz wavelets. *IEEE Transactions on Image Processing*, 23(2):898–908, February 2014.
- 13. Adrien Depeursinge, Antonio Foncubierta-Rodríguez, Dimitri Van De Ville, and Henning Müller. Three-dimensional solid texture analysis and retrieval in biomedical imaging: review and opportunities. *Medical Image Analysis*, 18(1):176–196, 2014.
- 14. Adrien Depeursinge, Camille Kurtz, Christopher F Beaulieu, Sandy Napel, and Daniel L Rubin. Predicting visual semantic descriptive terms from radiological image data: Preliminary results with liver lesions in CT. *IEEE Transactions on Medical Imaging*, 33(8):1–8, August 2014.
- 15. Adrien Depeursinge, Pedram Pad, Anne C Chin, Ann N Leung, Daniel L Rubin, Henning Müller, and Michael Unser. Optimized steerable wavelets for texture analysis of lung tissue in 3-D CT: classification of usual interstitial pneumonia. In *IEEE 12th International Symposium on Biomedical Imaging*, ISBI 2015, pages 403–406. IEEE, apr 2015.
- 16. Adrien Depeursinge, Zsuzsanna Püspöki, John-Paul Ward, and Michael Unser. Steerable wavelet machines (SWM): Learning moving frames for texture classification. *IEEE Transactions on Image Processing*, 26(4):1626–1636, 2017.
- 17. Adrien Depeursinge, Masahiro Yanagawa, Ann N Leung, and Daniel L Rubin. Predicting Adenocarcinoma Recurrence Using Computational Texture Models of Nodule Components in Lung CT. *Medical Physics*, 42:2054–2063, 2015.
- 18. Yashin Dicente Cid, Henning Müller, Alexandra Platon, Pierre-Alexandre Poletti, and Adrien Depeursinge. Locally-Oriented Wavelet Transforms for 3-D Solid Texture Classification. *IEEE Transactions on Image Processing*, submitted.
- 19. Qi Dou, Hao Chen, Yueming Jin, Lequan Yu, Jing Qin, and Pheng-Ann Heng. *3D Deeply Supervised*

 $\overline{)}$ 

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#### BIBLIOGRAPHY 99

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✐

*Network for Automatic Liver Segmentation from CT Volumes*, pages 149–157. Springer International Publishing, Cham, 2016.

- 20. Janis Fehr and Hans Burkhardt. 3D rotation invariant local binary patterns. In *19th International Conference on Pattern Recognition, 2008*, ICPR 2008, pages 1–4, December 2008.
- 21. William T Freeman and Edward H Adelson. The design and use of steerable filters. *IEEE Transactions on Pattern Analysis and Machine Intelligence*, 13(9):891–906, sep 1991.
- 22. Mary M Galloway. Texture analysis using gray level run lengths. *Computer Graphics and Image Processing*, 4(2):172–179, 1975.
- 23. Balaji Ganeshan, Elleny Panayiotou, Kate Burnand, Sabina Dizdarevic, and Kenneth A. Miles. Tumour heterogeneity in non-small cell lung carcinoma assessed by CT texture analysis: a potential marker of survival. *European Radiology*, 22(4):796–802, 2012.
- 24. Mehrdad J Gangeh, Ali Ghodsi, and Mohamed S Kamel. Dictionary Learning in Texture Classification. In *Proceedings of the 8th international conference on Image analysis and recognition - Volume Part I*, pages 335–343, 2011.
- 25. Diego Marcos Gonzalez, Michele Volpi, and Devis Tuia. Learning rotation invariant convolutional filters for texture classification. *CoRR*, abs/1604.0, 2016.
- 26. Ian Goodfellow, Aaron Courville, and Yoshua Bengio. Deep Learning. Book in preparation for MIT Press, 2015.
- 27. Zhenhua Guo, Lei Zhang, and David Zhang. A Completed Modeling of Local Binary Pattern Operator for Texture Classification. *IEEE Transactions on Image Processing*, 19(6):1657–1663, jun 2010.
- 28. Isabelle Guyon and Andre Elisseeff. An introduction to variable and feature selection. *Journal of Machine Learning Research*, 3:1157–1182, mar 2003.
- 29. Mohammad Haghighat, Saman Zonouz, and Mohamed Abdel-Mottaleb. CloudID: Trustworthy cloud-based and cross-enterprise biometric identification. *Expert Systems with Applications*, 42(21):7905–7916, 2015.
- 30. Robert M Haralick. Statistical and Structural Approaches to Texture. *Proceedings of the IEEE*, 67(5):786–804, may 1979.
- 31. Aapo Hyvarinen, Jarmo Hurri, and Patrik O. Hoyer. Energy Correlations and Topographic Organi- ¨ zation. In *Natural Image Statistics: A Probabilistic Approach to Early Computational Vision*, pages 239–261. Springer London, London, 2009.
- 32. Giovanni Jacovitti and Alessandro Neri. Multiresolution circular harmonic decomposition. *IEEE Transactions on Signal Processing*, 48(11):3242–3247, 2000.
- 33. Bela Julesz. Textons, the elements of texture perception, and their interactions. *Nature*, 290(5802):91–97, mar 1981.
- 34. James M Keller, Susan Chen, and Richard M Crownover. Texture description and segmentation through fractal geometry. *Computer Vision, Graphics, and Image Processing*, 45(2):150–166, 1989.
- 35. Alex Krizhevsky, Ilya Sutskever, and Geoffrey E. Hinton. ImageNet classification with deep convolutional neural networks. In F. Pereira, C. J. C. Burges, L. Bottou, and K. Q. Weinberger, editors, *Advances in Neural Information Processing Systems 25*, pages 1097–1105. Curran Associates, Inc., 2012.
- 36. Tai Sing Lee. Image representation using 2D Gabor wavelets. *IEEE Transactions on Pattern Analysis and Machine Intelligence*, 18(10):959–971, oct 1996.
- 37. Yuh-Jye Lee and Olvi L Mangasarian. SSVM: A smooth support vector machine for classification. *Computational optimization and applications*, 20(1):5–22, 2001.
- 38. Kun Liu, Henrik Skibbe, Thorsten Schmidt, Thomas Blein, Klaus Palme, Thomas Brox, and Olaf Ronneberger. Rotation-Invariant HOG Descriptors Using Fourier Analysis in Polar and Spherical Coordinates. *International Journal of Computer Vision*, 106(3):342–364, 2014.
- 39. L Liu, S Lao, P W Fieguth, Y Guo, X Wang, and M Pietikinen. Median Robust Extended Local Binary Pattern for Texture Classification. *IEEE Transactions on Image Processing*, 25(3):1368–1381, mar 2016.
- 40. David G Lowe. Distinctive Image Features from Scale–Invariant Keypoints. *International Journal of Computer Vision*, 60(2):91–110, 2004.

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#### 100 BIBLIOGRAPHY

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- 41. Julien Mairal, Francis Bach, J Ponce, Guillermo Sapiro, and Andrew Zisserman. Supervised dictionary learning. *Advances in Neural Information Processing Systems*, pages 1033–1040, 2008.
- 42. Benoit B Mandelbrot. *The Fractal Geometry of Nature*. W. H. Freeman and Company, 1977.
- 43. D Marr and E Hildreth. Theory of edge detection. *Proceedings of the Royal Society of London B: Biological Sciences*, 207(1167):187–217, feb 1980.
- 44. Fausto Milletari, Nassir Navab, and Seyed-Ahmad Ahmadi. V-Net: Fully Convolutional Neural Networks for Volumetric Medical Image Segmentation. *CoRR*, abs/1606.0, 2016.
- 45. Timo Ojala, Matti Pietikäinen, and Topi Mäenpää. Multiresolution gray–scale and rotation invariant texture classification with local binary patterns. *IEEE Transactions on Pattern Analysis and Machine Intelligence*, 24(7):971–987, jul 2002.
- 46. Maxime Oquab, Leon Bottou, Ivan Laptev, and Josef Sivic. Learning and Transferring Mid-level Image Representations Using Convolutional Neural Networks. In *2014 IEEE Conference on Computer Vision and Pattern Recognition*, pages 1717–1724, jun 2014.
- 47. Wumo Pan, T D Bui, and C Y Suen. *Rotation-Invariant Texture Classification Using Steerable Gabor Filter Bank*, pages 746–753. Springer Berlin Heidelberg, Berlin, Heidelberg, 2005.
- 48. A P Pentland. Fractal-Based Description of Natural Scenes. *IEEE Transactions on Pattern Analysis and Machine Intelligence*, PAMI-6(6):661–674, nov 1984.
- 49. B. Pesquet-Popescu and J. L. Vehel. Stochastic fractal models for image processing. *IEEE Signal Processing Magazine*, 19(5):48–62, sep 2002.
- 50. Maria Petrou and Pedro Garc´ıa Sevilla. *Non-Stationary Grey Texture Images*, pages 297–606. John Wiley & Sons, Ltd, 2006.
- 51. Javier Portilla and Eero P Simoncelli. A Parametric Texture Model Based on Joint Statistics of Complex Wavelet Coefficients. *International Journal of Computer Vision*, 40(1):49–70, 2000.
- 52. A G Ramakrishnan, S Kumar Raja, and H V Raghu Ram. Neural network-based segmentation of textures using Gabor features. In *Proceedings of the 12th IEEE Workshop on Neural Networks for Signal Processing*, pages 365–374, 2002.
- 53. Ali Sharif Razavian, Hossein Azizpour, Josephine Sullivan, and Stefan Carlsson. CNN features offthe-shelf: An astounding baseline for recognition. *IEEE Computer Society Conference on Computer Vision and Pattern Recognition Workshops*, pages 512–519, 2014.
- 54. Roberto Rigamonti and Vincent Lepetit. Accurate and Efficient Linear Structure Segmentation by Leveraging Ad Hoc Features with Learned Filters. In Nicholas Ayache, Hervé Delingette, Polina Golland, and Kensaku Mori, editors, *Medical Image Computing and Computer-Assisted Intervention MICCAI 2012*, volume 7510 of *Lecture Notes in Computer Science*, pages 189–197. Springer Berlin Heidelberg, 2012.
- 55. Olaf Ronneberger, Philipp Fischer, and Thomas Brox. U-Net: Convolutional Networks for Biomedical Image Segmentation. In Nassir Navab, Joachim Hornegger, William M Wells, and Alejandro F Frangi, editors, *Medical Image Computing and Computer-Assisted Intervention MICCAI 2015*, volume 9351 of *Lecture Notes in Computer Science*, pages 234–241. Springer International Publishing, 2015.
- 56. B Sahiner, Heang-Ping Chan, N Petrick, Datong Wei, M A Helvie, D D Adler, and M M Goodsitt. Classification of mass and normal breast tissue: a convolution neural network classifier with spatial domain and texture images. *IEEE Transactions on Medical Imaging*, 15(5):598–610, oct 1996.
- 57. N Sarkar and B B Chaudhuri. An efficient differential box-counting approach to compute fractal dimension of image. *IEEE Transactions on Systems, Man, and Cybernetics*, 24(1):115–120, jan 1994.
- 58. Paul Scovanner, Saad Ali, and Mubarak Shah. A 3-dimensional SIFT descriptor and its application to action recognition. In *Proceedings of the 15th ACM International Conference on Multimedia*, MM'07, pages 357–360, New York, NY, USA, 2007. ACM.
- 59. H Skibbe, M Reisert, T Schmidt, T Brox, O Ronneberger, and H Burkhardt. Fast Rotation Invariant 3D Feature Computation Utilizing Efficient Local Neighborhood Operators. *IEEE Transactions on Pattern Analysis and Machine Intelligence*, 34(8):1563–1575, aug 2012.
- 60. Guillaume Thibault, Bernard Fertil, Claire Navarro, Sandrine Pereira, Pierre Cau, Nicolas Levy, Jean Sequeira, and Jean-luc Mari. Texture Indexes and Gray Level Size Zone Matrix Application to Cell

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#### BIBLIOGRAPHY 101

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Nuclei Classification. *Pattern Recognition and Information Processing*, pages 140–145, 2009.

- 61. I Tosic and P Frossard. Dictionary Learning. *IEEE Signal Processing Magazine*, 28(2):27–38, mar 2011.
- 62. Michael Unser and Nicolas Chenouard. A unifying parametric framework for 2D steerable wavelet transforms. *SIAM Journal on Imaging Sciences*, 6(1):102–135, 2013.
- 63. Michael Unser, Nicolas Chenouard, and Dimitri Van De Ville. Steerable Pyramids and Tight Wavelet Frames in  $L_2(\mathbb{R}^d)$ . *IEEE Transactions on Image Processing*, 20(10):2705–2721, oct 2011.
- 64. Michael Unser and Dimitri Van De Ville. Wavelet steerability and the higher–order Riesz transform. *IEEE Transactions on Image Processing*, 19(3):636–652, March 2010.
- 65. Dimitri Van De Ville and Michael Unser. Complex wavelet bases, steerability, and the Marr-like pyramid. *IEEE Transactions on Image Processing*, 17(11):2063–2080, November 2008.
- 66. Manik Varma and Andrew Zisserman. A Statistical Approach to Texture Classification from Single Images. *International Journal of Computer Vision*, 62(1-2):61–81, 2005.
- 67. Andrea Vedaldi and Karel Lenc. MatConvNet: Convolutional Neural Networks for MATLAB. In *Proceedings of the 23rd ACM International Conference on Multimedia*, MM '15, pages 689–692, New York, NY, USA, 2015. ACM.
- 68. Jiazhuo Wang, John D MacKenzie, Rageshree Ramachandran, and Danny Z Chen. *A Deep Learning Approach for Semantic Segmentation in Histology Tissue Images*, pages 176–184. Springer International Publishing, Cham, 2016.
- 69. Y D Wang, Q Y Yan, and K F Li. Hand vein recognition based on multi-scale LBP and wavelet. In *2011 International Conference on Wavelet Analysis and Pattern Recognition*, pages 214–218, jul 2011.
- 70. J P Ward and M Unser. Harmonic Singular Integrals and Steerable Wavelets in *L*2(R*<sup>d</sup>*). *Applied and Computational Harmonic Analysis*, 36(2):183–197, mar 2014.
- 71. Dong-Hui Xu, Arati S Kurani, Jacob Furst, and Daniela S Raicu. Run–length encoding for volumetric texture. In *The 4th IASTED International Conference on Visualization, Imaging, and Image Processing – VIIP 2004*, Marbella, Spain, sep 2004.
- 72. Song-Chun Zhu, Cheng-En Guo, Yizhou Wang, and Zijian Xu. What are Textons? *International Journal of Computer Vision — Special Issue on Texture Analysis and Synthesis*, 62(1–2):121–143, apr 2005.