

Tumour Classification By Volumetric Image Analysis

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Abstract—This research review covers techniques available for binary classification of tumours in the human body considering practical implications of individual methods. An architecture is proposed in order to give direction to research and to allow consideration of stage interactions. Targeted at medical images the system follows standard image processing conventions and contains a sub-module that provides confidence information to domain expert. Only volumetric images processing is considered. Pre-processing and Feature extraction techniques are reviewed. Machine learning techniques for feature selection and classification are also considered.

The recommendation for population begins with anisotropic diffusion, for edge preservation smoothing, followed by edge detection and shape extraction. Sequential floating searching is used for feature selection when using a support vector machine for classification. Human supervised deployment is considered whether stand-alone or as an extension to existing medical review software.

I. INTRODUCTION

THIS research review covers the use of volumetric image analysis in medicine to accurately classify the presence of tumours in the human body. The problem is to be approached by considering the general cases so concentration is not applied to a specific area of the human body. Rather than trying to classify lung or brain tumours individually the goal is to consider what tumours have in common and given an entire image of a human body is it possible to classify tumours in any given section? The motivation behind this is to make the most of medical scanning. Dosages of radiation, cost and professionals time are all reasons to reduce the number of required patient scans. Thoroughly inspecting any data gathered is a constructive method of scan reduction. The objective is to provide a recommendation for a tool to be implemented to assist in medical practitioner diagnosis.

The architecture in Fig. 1 is the framework of the system to be recommended and is inspired by those discussed in [1]–[5]. A chain of stages leads from raw data gathered from the patient to a diagnosis from a classification algorithm. A confidence sub-block enables the domain expert to view the performance of the entire system without knowing the intricacies of operation; this requires processing to be explained with respect to their domain. In machine learning for medical applications a classification threshold has to be selected carefully as false negatives are more serious the false positive.

Raw data is captured from the patient and passed to system for pre-processing then feature extraction. Input patterns are produced dependent on a pre-determined list of known useful

tumour features. Dimensionality reduction is then used to frame the data as such to make the most of the classification algorithm.

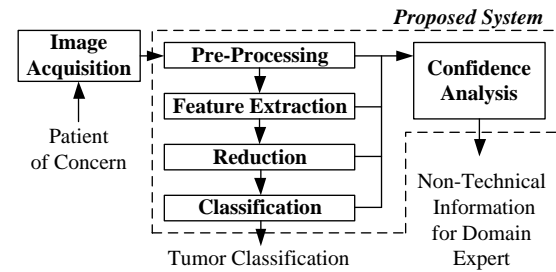


Fig. 1. Proposed system architecture.

II. IMAGE ACQUISITION

A standard medical X-ray is tuned to produce an image of human bone. The rays are attenuated by different materials in the body, more so by dense regions, and the image is gained from this attenuation effect [6]. This uses a single source and substrate to capture the image. Taking multiple images from many different angles gathers more information and a 3D image can be made by stitching these together. Fig. 2 describes the physical process of gathering data to be used in Computed Tomography (CT). A ring allows images to be captured from 360° in a single plane then by moving in the orthogonal plane, in Fig. 2 through the paper, a full volumetric image of the patient can be made.

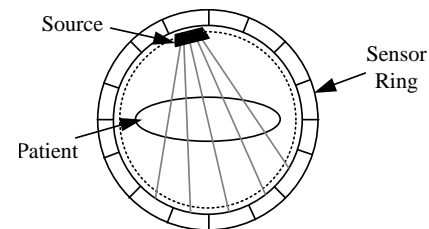


Fig. 2. Acquiring data from Computed Tomography. Adapted from [6].

Any method of non-invasive image capturing in 2D can therefore be expanded to work in 3D by using the CT principle. Different methods have trade-offs such as radiation dosages and resolution. Ultrasound is a method of image acquisition with the advantage of a small computational processing overhead allowing internal organs to be viewed in real-time which

is beneficial because it allows practitioners to view internal physical movement. It can be used for CT but higher resolution methods are usually favoured such as X-ray and Magnetic Resonance Imaging (MRI).

There are also other non-intrusive methods that do not use electromagnetic or ultrasonic waves. Hand palpitation is a common technique used by medical practitioners to gain an impression of abnormalities near the surface of the human body. Electromechanical apparatus employing the same technique can be used transfer the geometry of a growth to a volumetric image [7], [8]. Only part of the area under inspection is visible so the technique has obvious drawbacks.

Only high resolution grayscale sources will be considered in this review. This data is typically stored as a series of 2D images which are slices of a scan. An image viewer will build 3D images as and when required using these slices. The common medical standard for these images is DICOM which holds the slices, relative position and also the patient's details [9]. Viewing the data naturally proves problematic because it is 3D data shown in an isometric fashion on a 2D screen. Software provides interactive images as shown in Fig. 3 which allows the user to move three planes to gain an impression of the region.

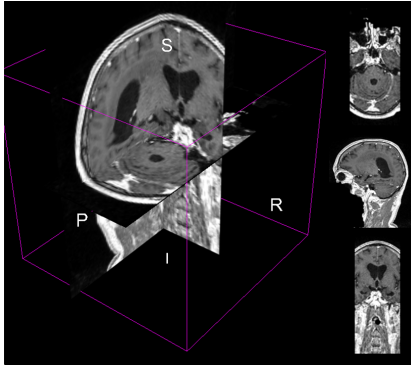


Fig. 3. An MRI scan of a healthy brain taken from [10] and rendered by [11].

III. TUMOUR FEATURES

There are hundreds of types of tumours that can occur in the human body but they are all caused by abnormalities in cell growth [12]. Tumours are dense areas of cells and can be either benign or malignant. Malignant tumours are considered cancers because they invade nearby normal tissue and spread therefore making them an area of concern. These are broken down in to three categories. *Carcinomas*, cells lining the body and internal organs, account for 90% of human caners. *Leukemias* and *Lymphomas*, occurring in the blood and lymph systems, account for 8% in humans. *Sarcomas* are solid regions of connective tissue which are rare.

Whether the tumour lines the host or grows independently the key to identification is to differentiate normal and abnormal cell density. Figure 4 contains examples of identified tumours in different organs. Tumours in brain (4(a)) and lung (4(c)) images are far simpler to identify than the liver image (4(c)). An image of a liver will contain other organs so these must be considered when trying to extract features.

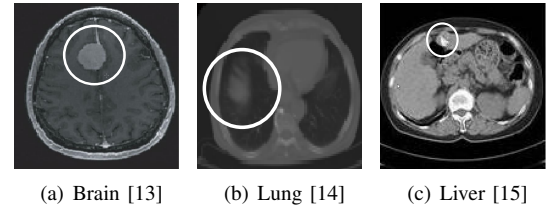


Fig. 4. Example MRI images containing tumours.

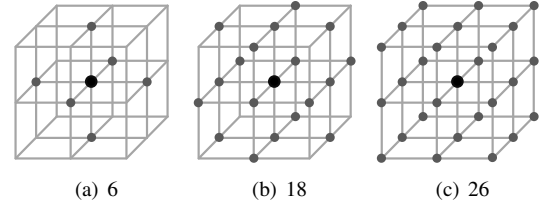


Fig. 5. Voxel Neighbourhoods [16].

IV. PRE-PROCESSING

The analogue of a pixel (picture element) in a 2D image is called a voxel (volumetric element) in a 3D image. This is exactly the same idea but with a extra dimension to form a cube [16]. They can be considered cubes but also points which may make visualisation and processing easier to understand. Fig. 5 describes how voxels are packed and the three different neighbourhoods that can be considered when inspecting a single voxel. Considering voxels to have larger neighbourhoods can increase the time complexity of some algorithms.

The raw data provided needs to be converted from the delivery format of the acquisition technology to a raw 3D matrix ready for generic processing. Watermarking, which may have been inserted for human review, should also be removed at this stage. This is trivial in theory but requires a database of artefacts or manual review to accomplish. There are also techniques, not considered in this report, which can be applied only to the slices of volumetric images therefore reducing computation overheads [17].

Pre-processing methods are used to enhance the image captured end ensure it is correctly formatted for further processing. This is considered as a mapping from an original image to a new image, $\mathbf{I} \mapsto \mathbf{I}'$, using only information from the original image.

A. Histogram Operations

An image histogram is a mapping from an image to a 2D plot of the frequency of voxel intensity. This is possible because voxels only have a discrete value of intensity within a fixed range. A histogram contains information on the type of material seen in the image, if it can be determined from intensity, and how much of that material is used to form the entire image. Image enhancement can be performed using point operations with information from the histogram. This is a very fast method of image enhancement as it is direct voxel-to-voxel mapping, as shown in Eq. (1). Normalisation stretches the histogram to cover all available values of intensity

therefore improving contrast. Equalisation attempts to flatten the histogram by distributing intensity [18].

$$I'(u, v, w) = \alpha(u, v, w)I(u, v, w) \quad (1)$$

B. Spatial Filtering

Spatial filters use the neighbourhood of voxels to create a new value for each voxel in an image. The original image is mapped through a filter kernel as described in Eq. (2). The filter kernel is a fixed dimension sub-image with a weighting on each voxel that is applied to every voxel in the original image. This is spatial convolution. If the filter kernel contains all ones it is known as a *box filter* and performs smoothing on the image to remove noise. A Gaussian filter performs better by placing a higher weighting on the central voxel and less weighting on the edge voxels inside the kernel [16]. The edge voxels of an image do not have a complete set of neighbours therefore performing spatial filtering will reduce the total size of the image in further processing.

$$I'(u, v, w) = \sum_{(i,j,k) \in R_H} I(u+i, v+j, w+k)H(i, j, k) \quad (2)$$

C. Frequency Filtering

The Fourier transform, as used spatially on 2D images, can be extended to 3D images. A Discrete Fourier Transform (DFT) for an N^3 voxel image is held in Eq. (3). The spatial filtering process performs convolution so once an image is in the frequency domain the same effects can be achieved using multiplication. It may prove more efficient to filter an image in the frequency domain rather than the spatial domain if the required filter kernel is large. Band-pass and notch filter operations can also be performed in the frequency domain. Eq. (4) provides the same result as Eq. (2) as described using Eq. (3).

$$F_{u,v,w} = \frac{1}{N^{\frac{3}{2}}} \sum_{x=-\frac{N}{2}}^{\frac{N}{2}-1} \sum_{y=-\frac{N}{2}}^{\frac{N}{2}-1} \sum_{z=-\frac{N}{2}}^{\frac{N}{2}-1} I_{x,y,z} e^{-j\frac{2\pi}{N}(ux+vy+wz)} \quad (3)$$

$$I' = \mathcal{F}^{-1}(\mathcal{F}(I) * \mathcal{F}(H)) = \mathcal{F}^{-1}(I * H) \quad (4)$$

D. Anisotropic Diffusion

Anisotropic diffusion is a powerful technique used to enhance images where it is possible to remove noise yet retain edges. The technique has been expanded from common 2D applications to 3D medical usage on MRI data [19]. The diffusion effect has the same outcome as a low pass filter with reasonably adjusted values for Δt and the decay constant. The spatial scalar k is a function of the image and must preserve edges by setting $k(u_{edge}, v_{edge}, w_{edge}, t) = 0$. Using Eq. (5) an iterative approach is provided in Eq. (6) where the original image is the value of I at time zero. Iteration monitoring and reasonable scalar values in the function k are required

to produce a good outcome. Edge detection methods to be used for k are discussed in section V-A.

$$\frac{\partial I}{\partial t} = k(I) \nabla^2 I, \quad k, I \in \mathbb{R}^4 \quad (5)$$

$$I_{t+\Delta t} = I_t + \Delta t k(I_t) \nabla^2 I_t \quad (6)$$

V. FEATURE EXTRACTION

High level feature extraction of volumetric shapes should be used on the input image. These high intensity regions may be organ walls or bones but should also include potential tumours.

A. Edge Detection

Edge detection can be performed using convolution with a kernel template as discussed in section IV-B. These templates are designed to create a new image contain voxel differences and can therefore be thought of as taking the derivative of the image. 2D templates are discussed in [18] and templates are the same for volumetric images. As all dimensions must be considered during differencing the 3D application is extended in the third dimension as shown in Fig 6. Smaller filter masks are also available to reduce computation but with diminished results.

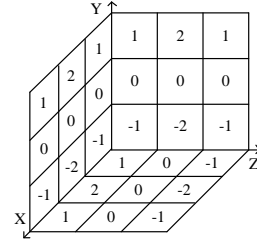


Fig. 6. 3D Sobel filter mask [16]

In edge critical processing better results can be produced with Canny edge detection [20]. A four stage process that uses Gaussian smoothing, the Sobel operator, non-maximal suppression followed by hysteresis thresholding. This reduces noise, minimises position difference in the mapping and eliminates multiple responses to an edge.

B. Generalised Hough Transform (GHT)

The GHT has been successfully applied to 3D images but as the shape of tumours are often unknown fitting a shape becomes redundant. Some tumours, Fig. 4(a), grow spherically so the ability to fit a sphere using the GHT may be a useful feature for classification.

C. Active Contours

The active contour model is a methods for flexible shape extraction. Expanding upon common 2D implementations active contours can be used to extract features from volumetric images. Eq. (7) contains the standard energy minimisation function [18], [21].

$$E = \int_0^1 E_{int}(\mathbf{v}) + E_{ext}(\mathbf{v}) + E_{con}(\mathbf{v}) ds, \quad \mathbf{v}(s) \in \mathbb{R}^3 \quad (7)$$

VI. FEATURE SELECTION

Commonly large datasets contain some features in the input space will be have no effect upon or even reduce the performance of a classifier. It is naive to use just the raw image data. Tumour features for extraction have been considered in section III which already removes some redundant information. Mapping input data into a lower dimensional space is possible and can improve performance but if ill-applied can remove key features. A brute-force approach would find the optimal combination of features but the input space is expected to be too large, 2^n , to be completely shattered within reasonable execution time.

A. Sequential Floating Search (SFS)

This heuristic algorithm is applied to tumour classification in [3]. It is possible to start from either the full or empty set of features then iteratively add and remove features to improve the performance of the classifier. A performance function is required for set comparison. Listing 1 contains the algorithm starting from an initial subset of no features and repeated until convergence.

```

1.  Z = {}, Y = {All}    // Setup
2.  Z = Z + Y(Best)      // Add the best feature
3.  X = Z - Z(Worst)     // Remove the worst
4.  if (f(X) > f(Z))
    Z = X
    goto 3              // Keep removing features
else
    goto 2              // Repeat

```

Listing 1. SFS Algorithm

B. Stochastic Approaches

A Random Mutation Hill Climber (RMHC) can move in the feature selection space from an initial starting point but will never reduce in performance. This does not handle a landscape with many local maxima well and may not produce a reasonable solution. Genetic algorithms are more computationally expensive as they require a population of solutions using crossover and mutation to converge on a suitable solution. Whether it outperforms a RMHC depends on the performance landscape but using a population of solutions is certainly more thorough. Simulated annealing can be used to move between solution neighbours subject over a fixed number of iterations. A cooling schedule is used to except negative moves in performance with a probability the is a function of the cooling variable.

C. Principle Component Analysis (PCA)

PCA is a common technique used for dimensionality reduction by ranking dimensions in order of variance [22]. This will rank the features so that only the first L are passed on for further processing. The other dimensions can simply be removed but if the effect of a low variance dimension is important to classification this can reduce performance. Eq. (8) shows how the original input pattern is mapped to a reduce pattern by subtracting the mean of all patterns and multiplying

with the projection matrix. The projection matrix is built by finding the covariance matrix of the input patterns then ranking the eigenvectors by the eigenvalues and deleting as required. Performance feedback from training can be used to select appropriate L value.

$$z = P(x - \mu) \quad (8)$$

VII. CLASSIFICATION

Classification must differentiate between extracted geometric shapes that are normal and those which may potentially be tumours. Tumours will also be attached to normal internal body parts so the classifier must be able to handle any abnormalities in organs. Supervised and unsupervised techniques for classification are considered even though a datasets have not yet been sourced.

A. Clustering

B. Artificial Neural Networks (ANNs)

ANNs contain summation nodes where data is propagated through weighted connections resulting in a final output. As a supervised technique weights are trained by propagating the error backwards through the network. Multilayer nets can provided non-linear separation functions and increasing the number of layers only increases possible function complexity.

C. Support Vector Machines (SVMs)

An SVM classifier builds a separation margin supported by the closest data points the drawing a separation plane from the middle of this margin. A linear maximum-margin hyperplane is produce but non-linear data can also be separated by using the kernel trick [23]. This is a mapping of the input space into a higher dimensional space so that it becomes linearly separable. SVMs have been successfully applied to breast cancer diagnosis in [24] The approach uses a weighted combination of polynomial and radial basis functions to improve the generalisation performance.

D. Boosting

A weak learner for binary classification can be considered any classifier with an error rate less than 0.5. It is possible to create a strong learner, having a much smaller error rate, by combing the weighted sum of different weak learners. The algorithm can be considered a hybrid of feature selection and classification. AdaBoost is the most common implementation of a boosting algorithm [25] and has been successfully applied to brain tumour classification in [26].

VIII. CONCLUSIONS

It is clear that this tool can't be used as a hard and fast method of tumour diagnosis but instead as a valuable tool for image reviewers. Software packages designed for medical image review, such as [11], have bidirectional interfaces to custom extensions. Implementing this system as an extension would remove many overheads required in data handling but reduce the scope for optimisation. The feedback from the entire process would be a checklist which the user must manually review and waive if incorrectly identified.

REFERENCES

- [1] S. Ahmed, K. Iftekharuddin, and A. Vossough, "Efficacy of texture, shape, and intensity feature fusion for posterior-fossa tumor segmentation in mri," *Info Tech in BioMed, IEEE Trans on*, vol. 15, no. 2, pp. 206–213, March 2011.
- [2] V. Kumar, J. Sachdeva, I. Gupta, N. Khandelwal, and C. Ahuja, "Classification of brain tumors using pca-ann," in *Info and Comms Tech (WICT), 2011 World Congress on*, Dec 2011, pp. 1079–1083.
- [3] H.-L. Bu, G.-Z. Li, X.-Q. Zeng, J. Yang, and M. Yang, "Feature selection and partial least squares based dimension reduction for tumor classification," in *Bioinformatics and Bioengineering, 2007. BIBE 2007. Proceedings of the 7th IEEE International Conference on*, Oct 2007, pp. 967–973.
- [4] J. Sachdeva, V. Kumar, I. Gupta, N. Khandelwal, and C. Ahuja, "Multiclass brain tumor classification using ga-svm," in *Developments in E-systems Eng. (DeSE), 2011*, Dec 2011, pp. 182–187.
- [5] W. Kostis, A. Reeves, D. Yankelevitz, and C. Henschke, "Three-dimensional segmentation and growth-rate estimation of small pulmonary nodules in helical ct images," *Med Imaging, IEEE Trans on*, vol. 22, no. 10, pp. 1259–1274, Oct 2003.
- [6] K. Najarian and R. Slinger, *Biomedical Signal and Image Processing*, 1st ed. Taylor and Francis Group, 2006.
- [7] H. L. Liu, O. Elhage, P. Dasgupta, B. Challacombe, D. Murphy, L. Seneviratne, and K. Althoefer, "A haptic probe for soft tissue abnormality identification during minimally invasive surgery," in *Reconfigurable Mechanisms and Robots, 2009. ReMAR 2009. ASME/IFToMM International Con. on*, June 2009, pp. 417–422.
- [8] P. S. Wellman and R. D. Howe, "Modeling probe and tissue interaction for tumor feature extraction," *ASME-PUBLICATIONS-BED*, vol. 35, pp. 237–238, 1997.
- [9] NEMA. (2011) DICOM. [Visited] 13/03/14.
- [10] FNLCR. The cancer imaging archive. [Visited] 16/03/14. [Online]. Available: <http://cancerimagingarchive.net/>
- [11] 3DSlicer. [Visited] 13/03/14. [Online]. Available: <http://www.slicer.org/>
- [12] G. Cooper, *Elements of Human Cancer*, ser. Biology Series. Jones and Bartlett Publishers, 1992.
- [13] R. Islam, A. Mamun, M. Bhuiyan, and S. Rahman, "Segmentation and 3d visualization of volumetric image for detection of tumor in cancerous brain," in *Elec Comp Eng (ICECE), 2012 7th Int. Con.*, Dec 2012, pp. 863–866.
- [14] A. Amutha and R. Wahidabanu, "Lung tumor detection and diagnosis in ct scan images," in *Comms and Sig. Proc. (ICCSP), 2013 Int. Con.*, April 2013, pp. 1108–1112.
- [15] Y. Chen, R. Xu, S.-Y. Tang, S. Morikawa, and Y. Kurumi, "Non-rigid mr-ct image registration for mr-guided liver cancer surgery," in *Complex Med Eng, 2007. CME 2007. IEEE/ICME Int. Con. on*, May 2007, pp. 1756–1760.
- [16] G. Lohmann, *Volumetric Image Analysis*, 1st ed. Wiley, 1998.
- [17] G. Harauz and M. van Heel, "Exact filters for general geometry three dimensional reconstruction," in *Proceedings of the IEEE Computer Vision and Pattern Recognition Conf*, vol. 73, 1986, pp. 146–156.
- [18] M. Nixon and A. Aguado, *Feature Extraction and Image Processing*, 1st ed. Newnes, 2002.
- [19] B. Nakhjavanlo, T. Ellis, P. H. Soan, and J. Dehmeshki, "3d medical image segmentation using level set models and anisotropic diffusion," in *Signal-Image Technology and Internet-Based Systems (SITIS), 2011 Seventh International Conference on*, Nov 2011, pp. 403–408.
- [20] J. Canny, "A computational approach to edge detection," *Pattern Analysis and Machine Intelligence, IEEE Transactions on*, vol. PAMI-8, no. 6, pp. 679–698, Nov 1986.
- [21] A. Skalski, J. Lagwa, P. Kedzierawski, T. Zielinski, and T. Kuszewski, "Automatic prostate segmentation in mr images based on 3d active contours with shape constraints," in *Signal Processing: Algorithms, Architectures, Arrangements, and Applications (SPA), 2013*, Sept 2013, pp. 246–249.
- [22] K. Pearson, "On lines and planes of closest fit to systems of points in space," *Phil. Mag.*, vol. 6, pp. 559–572, 1901.
- [23] C. Cortes and V. Vapnik, "Support-vector networks," *Machine learning*, vol. 20, no. 3, pp. 273–297, 1995.
- [24] X. Yang, H. Peng, and M. Shi, "Svm with multiple kernels based on manifold learning for breast cancer diagnosis," in *Information and Automation (ICIA), 2013 IEEE International Conference on*, Aug 2013, pp. 396–399.
- [25] Y. Freund and R. E. Schapire, "A short introduction to boosting," vol. 14, no. 5, Sep 1999, pp. 771–780.
- [26] A. Islam, S. Reza, and K. Iftekharuddin, "Multifractal texture estimation for detection and segmentation of brain tumors," *Biomedical Engineering, IEEE Transactions on*, vol. 60, no. 11, pp. 3204–3215, Nov 2013.