

Automated Registration of Ultrasound with *CT* Images: Application to Computer Assisted Prostate Radiotherapy and Orthopedics

Gelu Ionescu, PhD., Stéphane Lavallée, PhD., Jacques Demongeot, MD., PhD.

TIMC Laboratory, University Joseph Fourier, Grenoble France
gelu.ionescu@imag.fr

Abstract. Ultrasound (*US*) imaging could potentially play a major role in the field of Computer Assisted Surgery (CAS). For doctors and surgeons to make full use of CAS tools in planning and executing surgical operations, they also need user-friendly automatic software based on fast, precise and reliable registration algorithms. The main goal of this paper is to take advantage of the segmentation/registration duality to extract the relevant information from *US* images. This information will allow the precise and automatic registration of anatomical structures contained in the pre-operative model and of intra-operative data contained in *US* images. The result of registration will be further used to guide a computer-assisted intervention such as orthopedics or radiotherapy.

1 Introduction

Among the various imaging techniques available, *2D* echography is becoming more and more important, both for the diagnostic purposes and registration in CAS applications. In this paper, we will consider the latter case, with the specific instance of Computer Assisted Orthopedics Applications [1,2] and patient positioning in prostate radiotherapy [3]. In spite of the problems inherent to *US* imaging such as low signal-to-noise ratio and the fact that the images depend on the angle and the texture of the target anatomy, this modality has become very popular among practitioners mainly because of its safety, low cost and non-invasiveness.

Since external radiotherapy makes use of the properties of *X – rays* to destroy live tissue, the precise localization of the tumor is extremely important in order to avoid the destruction of neighboring healthy tissues. As described in details in [4], in the context of isocentric technique applied to radiotherapy, the procedure for repositioning must ensure that the center of the tumor localized by the practitioner in the pre-operative model, coincides with the center of rotation of the radiation device. For the intra-operative data acquisition, we are interested particularly in the transpubic way to realize the *US* examination of the prostate. The goal of the registration is to realize an automatic orientation of the external *X – ray* beams according to a predetermined dosimetry plan. The results presented here were obtained on patients. In orthopedic applications, a

high precision is also required in order to respect the planning established by the surgeon. Our aim is either to place a screw in a pedicle vertebra [2], or to insert percutaneously screws in the sacro-iliac bone [1,5]. In both cases, the goal is to realize a minimally invasive operation. To demonstrate the feasibility of the method, we worked, for the first application, on a plastic vertebra immersed in water, for the second application tests were performed on a cadaver.

In this paper, we present an automatic, reliable and precise method that can be used in soft tissues (e.g. abdominal cavity) as well as in hard tissues (e.g. bone, cartilage). The medical imagings we focus mainly on, are computed tomography (*CT*) and *US* imaging. The former is used to build the pre-operative 3D model and the latter to acquire the 3D intra-operative data volume.

2 Related Work

The general problem of registration between a set of 3D data points and a 3D model was treated in [6,7]. In this paper, we consider a 3D model made from a set of *CT* slices. This approach has been discussed in [2,5] for various applications. The methods known in the literature is based on the segmentation of the surface S of an anatomical structure on *CT* images. This is followed by a manual segmentation of pieces of contours corresponding to this same structure on *US* images. Each *US* image being located in the intra-operative reference frame by using an external optical localization device, one obtains an image of points belonging to the surface of the reference structure. The rigid registration of 3D points with the surface S starting from an initial position is described in various papers [7,8,9]. Unfortunately none of the existing segmentation procedure of *US* images is automatic, precise and robust. Here we propose a method featuring these characteristics [10].

Grey-Level Approach The most obvious method, as often used in registration, would be to register directly *CT* images with *US* images without going through segmentation. For images originating from the same modality, one could use methods based on maximization of correlation, confinement tree [11]. For images of different nature, optimization based on mutual information could be used [12]. For a given initial registration T_0 , for each *US* image in a given position, one can calculate the slice corresponding to the echographical plane in the volume of *CT* images but the calculation of reformatted slices is very time consuming [10]. For this reason an approach based on the use of contours seemed preferable.

Contour Approach As a starting point, the 3D model is assumed to be both segmented and labeled. To achieve this it is of course possible to use semi-interactive procedures since the model is constructed before the operation. In general, one segments contours on 2D slices. These are then interpolated to create a homogeneous 3D surface using the shape-based interpolation algorithm described in [13,10]. Although many registration/3D labeling procedures are described in the literature, those based on elastic registration of a model (Atlas

like) seem the most promising ones [14]. The segmentation/registration duality was discussed previously in different context. Hamadeh [15] proposed to register the 3D model of a segmented vertebra on *CT* images with intra-operative *X-ray* of the same vertebra. Mangin [16] proposed a two-step registration of Positron Emission Tomography and *CT* images of the brain while recently Gerlaud [17] proposed a progressive approach using Atlas-type models to segment the Magnetic Resonance images of the brain. Our approach, as in the above cited works, is based on the segmentation/registration duality, its specificities are that we rely on *US* imaging and that we propose an original mechanism for segmentation, guided by the model. As in the method based on gray-levels, we propose to generate pseudo *US* images, however, we strictly limit ourselves to the contours of structures segmented in the model. This is equivalent to considering only the specular reflection and to neglect all other phenomena. This approach is detailed in the following sections.

3 Methods

The method we have proposed consists of three stages, the first consists of a low-level segmentation of a sequence of *US* images. The second stage exploits the *segmentation/registration* duality to extract relevant contours obtained in the previous step. The last step consists of the final rigid registration of these relevant contours and the pre-operative model.

3.1 Intra-operative Data Acquisition

Images are taken from an *US* imager. The *US* probe is localized in 3D space by an optical localizer. The *US* image is calibrated according to a technique described in [10]. In this way, the position of an image pixel is known in 3D space with a precision in the range of localizer accuracy (about 1 mm).

3.2 Low-Level Segmentation

By low-level segmentation, we mean classical image segmentation techniques based mainly on linear filtering or mathematical morphology. At the end of this stage, the images are generally over-segmented (presence of false positives) [18] and the contours are not labeled. In this work, we choose the watershed as the segmentation technique. The main advantages of watershed segmentation relative to methods based on filtering [19] are the following: the watershed always contains some segments placed correctly on the real contours, the resulting contours are closed, detection of multiple junctions is very insensitive to noise and the algorithm is based on the modulus of the gradient $\|\nabla I\|$ of the image (the direction of the gradient is ignored).

To eliminate undesirable effects produced by the speckle on the complexity of the watershed, the original image is filtered by a recursive Canny-Deriche filter [20]. The width of this filter is chosen in an empirical way but it remains

unchanged for the processing of the whole sequence of images. Furthermore, we observed that the filter width does not change significantly from one patient to the next. For the test-patient used for the prostatic application, the parameter for the Canny-Deriche filter is $\alpha = 0.2$. The result of this first step presented in Fig. 1.e shows two false watershed segments going through the bladder/prostate structure and the right iliac bone. One notes immediately that the prostate is well isolated which demonstrates that the use of watershed is completely realistic. At this stage, the processing is specific to the structure that needs to be detected.

3.3 High-Level Segmentation

At this stage, we assume the availability of a set of M initial attitudes $T_i (i = 1, M)$, relating the echographical data and the *CT* model. Each initial attitude T_i is applied to each of the N echo slices. One then obtains M possible intersections between each echo plane and the dense cluster of points representing the surface of the model. In order to increase the resolution for the exploration of the volume of the model, to each initial attitude T_i , one associates six additional transformations $\Delta T_j (j = 0, 6)$ which slightly displace the image plane as presented in Fig. 2.a. In this scenario, we voluntarily neglected the displacements inside the image plane since these could be compensated by the $2D/2D$ elastic registration described in this section.

For each combination $(T_i, \Delta T_j)$, we keep the model points which lie within the thickness of the *US* beam ($\simeq 2\text{ mm}$). We then perform an elastic multi-level $2D/2D$ registration between the simulated echo slice and the real echo slice. As described hereinafter, we minimize an expression of the form:

$$\varepsilon = \sum_{i=1}^N \text{dist}_{2D} (Q_j, T_p(P_i))_{i=1,M} \quad (1)$$

where P_i is the set of N simulated echo points, Q_j is the set of M real echo points and T_p is the elastic $2D$ transformation relative to the parameter vector p we are looking for.

In order to give to the registration process the necessary elasticity, we associate to the image a regular network, the nodes of which can be individually adjusted. A displacement vector $(V_x, V_y)^\tau$ is associated to each node as presented in Fig. 2.b. The global transformation T_p we are looking for is parameterized by the vector p :

$$p = \left(\alpha, t_x, t_y; (V_x, V_y)_{0,0}^\tau, \dots, (V_x, V_y)_{i,j}^\tau, \dots, (V_x, V_y)_{I_x, I_y}^\tau \right)^\tau \quad (2)$$

where τ means transposition, (I_x, I_y) is the network dimension, $(\alpha, t_x, t_y)^\tau$ is the purely rigid component and $\left((V_x, V_y)_{0,0}^\tau, \dots, (V_x, V_y)_{i,j}^\tau, \dots, (V_x, V_y)_{I_x, I_y}^\tau \right)^\tau$ is the purely elastic component of the transformation T_p . To determine the vector p , we have developed a nonlinear iterative optimization algorithm based on the

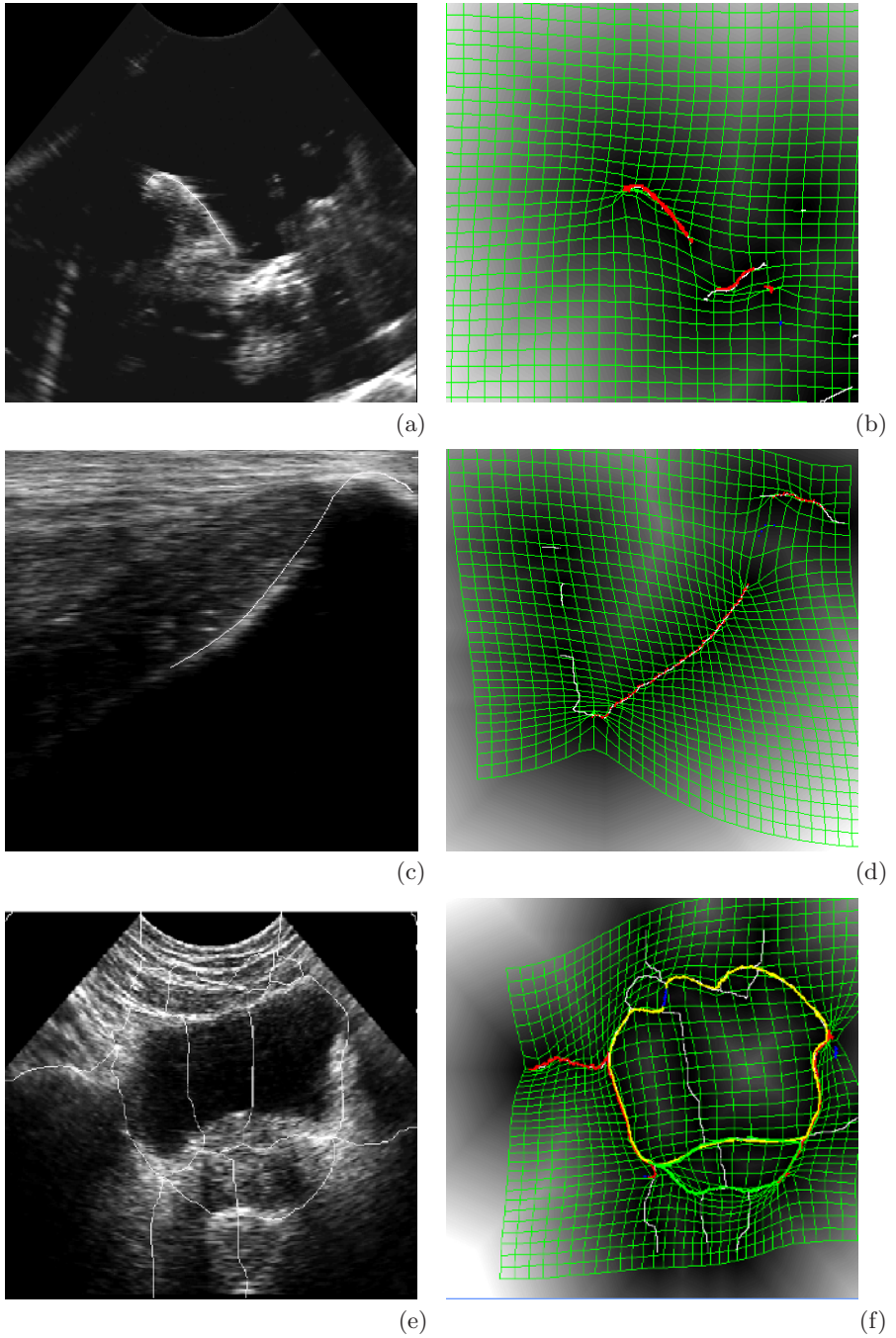
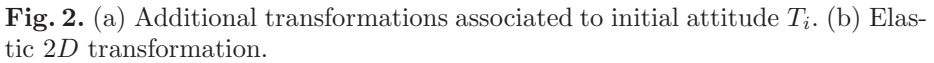


Fig. 1. Typical *US* images (left) and 2D/2D elastic registration (right). (a-b) Vertebra. (c-d) Sacro-iliac region. (e-f) Bladder and prostate.


$$\begin{aligned} \varepsilon = & \frac{1}{N} \sum_{i=1}^N \|Q_j - T_p(P_i)\|^2 + \rho_1 \frac{1}{I_x \times I_y} \sum_{(i,j)=[0,0]}^{[I_x, I_y]} \|V_{i,j}\|^2 + \\ & \rho_2 \frac{1}{I_x \times I_y} \left(\sum_{(i,j)=[0,0]}^{(I_x, I_y)} \|V_{i+1,j} - V_{i,j}\|^2 + \sum_{(i,j)=[0,0]}^{[I_x, I_y)} \|V_{i,j,1} - V_{i,j}\|^2 \right) \end{aligned} \quad (3)$$

To speed up the computation of distances, we use the 2D chamfer distance map for the set of real points Q_j . The minimization method used is based on the calculation of gradient and uses the following iterative mechanism:

$$\delta p_k = -\lambda \nabla \varepsilon_k \quad (5)$$

Convergence Criteria The criteria for stopping the iterative procedure are related to the energy and the gradient:

1. $\frac{\varepsilon_{t+1}^2 - \varepsilon_t^2}{\varepsilon_t^2} \leq T_1$; the relative energy gain between two successive successful iterations must be below a reasonable threshold T_1 .

2. $N_{iterations}$; after $N_{iterations}$ successive unsuccessful iterations.
3. (I_x, I_y) ; after the convergence using the previously defined criteria, the network resolution can be increased and the minimization starts again.

Retaining the Best $(T_i, \Delta T_j)$ Combination In the case of hard tissues, the choice of the best $(T_i, \Delta T_j)$ combination relies on a likelihood principle. Consequently, we construct an objective discriminating function based on the first term of (3) and on the number of points P_i of the model. Among many possibilities, the solution retained is based on the Student test t in order to compare variances. We have adopted the solution proposed in [21] where the original data are transformed in the absolute values of the deviation of each distance relative to the average value using the relation:

$$t = (\bar{x}_1 - \bar{x}_2)s\sqrt{\frac{1}{n_1} - \frac{1}{n_2}} \quad (6)$$

$$s = ((n_1 - 1)s_1^2 + (n_2 - 1)s_2^2) / (n_1 + n_2 - 2) \quad (7)$$

where $n_{1,2}$ is the sample size, $\bar{x}_{1,2}$ is the sample mean value, $s_{1,2}$ is the sample variance and s is the global variance. The value $t_{k,l}$ obtained from expression (6) is used as a dissimilarity measure for the couple $[(T, \Delta T)_k, (T, \Delta T)_l]$. A symmetric matrix $T = [t_{k,l}]; k, l \in [1, N \cdot M]$ is then constructed. An objective function $f_t(l) = \sum_{k=1}^{N \cdot M} t_{k,l}$ is obtained by summing over the columns of the matrix T . The decision mechanism is based on minimizing $f_t(l)$. Consequently, the optimal $(T, \Delta T)_{opt}$ combination corresponds to the relation $(f_t)_{opt} = \min_{l=1, N \cdot M} f_t(l)$.

In the case of soft tissues, one cannot defend the use of maximum likelihood as an objective discriminatory function. For instance it is very easy to register elastically two transversal slices through the bladder. Taking into account the preceding observation, we propose a solution adapted to all soft tissues based on two criteria, a quantitative one followed by a qualitative one: 1. Establish a measure that determines how well the points of the model fit the intra-operative data. To achieve this, we compute a distance map based on the new coordinates of the deformed model points taken after the $2D/2D$ elastic registration. From this distance map, we obtain the distance of each intra-operative data point relative to the model. The average value of all these distances, $\bar{d}_{i,j}$, defines the objective measure characterizing the $(T_i, \Delta T_j)$ combination. The best combination being the one with an uniform distribution of the points of the deformed model as compared with intra-operative data. With a minimization procedure one obtains $(T, \Delta T)_{opt} = \min_{l=1, N \cdot M} (\bar{d}_{i,j})_l$. 2. In problematic situations when none of the $(T_i, \Delta T_j)$ combinations represent a realistic situation (the model points are not well-distributed) this image will be marked and eliminated from the final $3D/3D$ registration process. The decision criteria are the following: the model points must cover at least 50 % of the data and in the case of the prostate, this area must necessarily cover the base of the bladder. After the choice of the best $(T, \Delta T)_{opt}$ combination, another step of $2D/2D$ elastic registration will be necessary. This time, intra-operative data are represented by the watershed

while pre-operative model is represented by the set of anatomic structure of the region of interest (i.e. the bone, the bladder and the prostate). The results are presented in Fig. 1.

3.4 Rigid Registration

Choice of Segments For this step, we had to choose between two possibilities: isolate all segments of the watershed pointed by the model or retain the new coordinates of the points of the model. To show that both possibilities are realistic, we used the first one in orthopedic applications and the second one in radiotherapy applications.

A final *3D/3D* registration step is necessary to match the points previously extracted and labeled on *US* images with the surface of the *CT* model. For that purpose, one could use elastic registration [22], but both the prostate (for radiotherapy) and the bones (for orthopedics) are considered as rigid bodies. Therefore, standard *3D* surface registration techniques were used to estimate the transformation between *CT* model and the patient coordinate system in the local region of interest [23].

4 Results

In order to validate our technique, we compared the final results obtained after the rigid *3D/3D* registration with either those obtained from a manual segmentation of the prostate or, in the case of the orthopedic application, with manual digitalization of the bone surface with a finger probe localized in *3D* space. All numerical results are presented in the following table where we display the errors obtained both on rotations $\varepsilon_{\max}^{\alpha}$ expressed in degrees and on translations ε_{\max}^t expressed in *mm*. In the last column, the same results are presented in terms of Rodriguez's rotation and translation.

| | $\varepsilon_{\max}^{\alpha}[\text{deg}]$ | $\varepsilon_{\max}^t[\text{mm}]$ | <i>Rodriguez</i> |
|----------|---|-----------------------------------|------------------|
| prostate | (-1.23, 1.30, 0.75) | (0.19, -0.67, -2.58) | (2.67°, 2.07mm) |
| vertebra | (1.90, 2.05, -0.98) | (-0.15, -0.52, 0.87) | (1.23°, 1.70mm) |
| bassin | (-2.93, 1.13, -1.83) | (-0.39, 0.12, -1.02) | (1.14°, 1.91mm) |

In Fig. 3, we present the repositioning of a typical *US* image in *3D* space of the *CT* per-operative model. Fig. 3.a displays the superposition of the model on the spinal process of the vertebra, Fig. 3.b presents a superposition with the *US* image representing the sacrum and Fig. 3.c displays an *US* image of the bladder and the prostate.

5 Conclusion

In this paper, we have presented automatic and robust algorithms necessary for registration between a *3D CT* model and a set of *2.5D US* images. Essentially, these algorithms use methods of low-level segmentation as described in

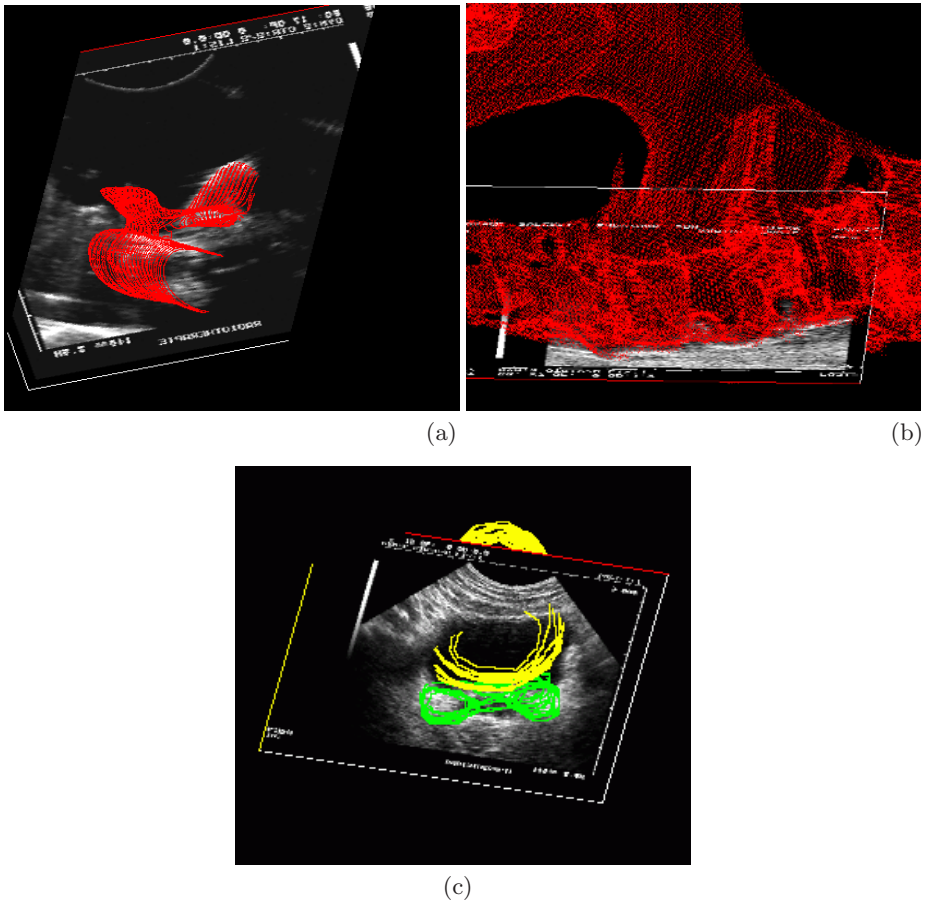


Fig. 3. Results: (a) Vertebra registration. (b) Sacrum registration. (c) Bladder and prostate registration

paragraph §3.2 and high-level segmentation by labeling as described in §3.3. To emphasize the generality of our approach, each application has been analyzed in the following framework: description of clinical objectives, presentation of specific pre-processing, labeling by elastic $2D/2D$ registration, choice of segments and presentation of results. In all this work, we paid special attention to the precision and we could demonstrate that the maximum errors are about 2 mm and 2^0 which is compatible with most of applications. While the orthopedic application chosen clearly deals with high-risk regions, in the case of the prostate, the risk is to irradiate healthy organs. Considering the important risks inherent to an intervention on the spine or in the sacral region, we have tested our method *in vitro*. The preliminary results obtained in the automatization of the segmentation process are very encouraging. It is clear that the result of a minimization

procedure generally used in matching depends strongly on the precision of the initial attitude T_0 , this also affects the global computing time. To improve on this, we hope to find a procedure that estimates an initial attitude T_0 as close as possible to the final attitude for each application. We plan also to correct the *US* images, in term of the velocity of *US*. This will influence the precision of the results specially in the radiotherapy applications. This method is generic and it could be easily applied to other organs such as liver and kidney.

References

1. Tonetti J. et al, Percutaneous iliosacral screw placement using image guided techniques, *CORR special issue*, pp. 103-110, 1998. 768, 769
2. Merloz P. et al, Pedicle screw placement using image guided techniques, *CORR special issue*, pp. 39-48, 1998. 768, 769
3. Troccaz J. et al, Conformal external radiotherapy pf prostatic carcinoma: requirements and experimental results, *Radiother. Oncol.*, no. 29, pp. 176-183, 1993. 768
4. Vassal P., Fusion d'images multi-modales pour la radioth. conformationnelle: appli. au repos. du patient, *Ph.D. Thesis TIMC-IMAG, Grenoble University*, 1998. 768
5. Carrat L. et al, Treatment of pelvic ring fractures: percutaneous computer assisted iliosacral screwing, *MICCAI'98-Proceedings*, pp. 84-91, 1998. 769
6. Lavallée S. et al, Computer integrated surgery and therapy, *C. Roux and J.L. Coatrieux ed., Contemp. Persp. in 3D Biomed. Im. IOS Press*, pp. 239-310, 1997. 769
7. Maintz A. and Viergever M.A., A survey of medical image registration, *Medical Image Analysis*, vol. 2, no. 1, pp. 1-36, 1998. 769
8. Pelizzari C.A. et al, Accurate 3D registration of CT, PET and/or MR images of brain, *L. Comput. Assist. Tomo.*, no. 13, pp. 20-26, 1989. 769
9. Lavallée S. et al, Matching 3D smooth surf. with their 2D proj. using 3D dist. maps, *SPIE vol. 1570 Geom. Models in Computer Vision - San Diego*, pp 322-336, 1991. 769
10. Ionescu G., Segm. et recalage d'images échographiques par utilisation de connaissances physio. et morpho., *Ph.D. Thesis TIMC-IMAG, Grenoble University*, 1998. 769, 770
11. Mattes J. and Demongeot J., Dynamic confinement, classification and imaging, *New techn. of classific., Lecture Notes in Comput. Sci., Springer Verlag*, 1998. 769
12. Wells W.M. et al, Multimodal volume registration by minimisation of mutual information, *Medical Robotics and CAS Wiley New York*, pp 55-62, 1995. 769
13. Grevera G.J. and Udupa J.K., Shape-based interpolation of multidimensional grey-level images, *IEEE Trans. Medical Imaging*, vol. 15, no. 6, pp. 881-891, Dec. 1996. 769
14. Bittar E., Modèles déformables surfaciques, implicites et volumiques, pour l'imagerie médicale, *Ph.D. Thesis TIMC-IMAG, Grenoble University*, 1998. 770
15. Hamadeh A., Une approche unifiée pour la segm. el la mise en corresp. 3D/2D d'images multi-modales, *Ph.D. Thesis TIMC-IMAG, Grenoble University*, 1997. 770
16. Mangin J., Mise en correspondance d'images médicales 3D multi-modalités pour la corrélation anatomo-fonctionnelle cérébrale, *Ph.D. Thesis ENST Paris*, 1995. 770

17. Géraud T., Segmentation des structures internes du cerveau en IRM 3D, *Ph.D. Thesis TLMCOM Paris*, 1998. 770
18. Maes F., Segmentation and registration of multimodal medical images, *Ph.D. Thesis ISBN 90-5682-135-0 D/1998/7515/37*, 1998. 770
19. Najman L. and Schmitt M., Geodesic saliency of watershed contours and hierarchical segmentation, *IEEE Trans. PAMI*, vol. 18, no. 12, pp. 1163-1173, Dec. 1996. 770
20. Deriche R., Fast algorithms for low-level vision, *IEEE Trans. PAMI*, vol. 12, no. 1, pp. 78-87, Jan. 1990. 770
21. Manly B., Statistical methods, *Chapman and Hall*, 1989. 774
22. Szeliski R. et al, Matching 3-D anatomical surfaces with non-rigid deformations using octree-splines, *Intern. Journ. of Comp. Vis.*, 18(2), pp. 171-186, 1996. 775
23. Lavallée S. et al, Recovering the pos. and orient. of free-form obj. from image contours using 3D dist. maps, *IEEE Trans. PAMI*, vol. 17, no. 4, pp. 378-390, 1995. 775