Measurement of Nuchal Translucency Thickness in First Trimester Ultrasound Foetal Images **Using Markov Random Field**

Journal of Pharmacology and Pharmacotherapeutics 15(1) 46-53, 2024 © The Author(s) 2024 Article reuse guidelines: in.sagepub.com/journals-permissions-india DOI: 10.1177/0976500X241232530 journals.sagepub.com/home/pha



Kalyani Chaudhari¹ and Shruti Oza²

Abstract

Background: The first-trimester ultrasound assessment of nuchal translucency (NT) thickness has lately been recommended as the most helpful sign in early screening for prenatal chromosomal disorders. Increased foetal NT thickness between II and 13+6 weeks of gestation is a frequent phenotypic manifestation of chromosomal abnormalities as well as a variety of foetal deformities and genetic disorders.

Purpose: At the moment, clinicians conduct the measurement manually. The measurement may take a long time to complete, requires highly competent operators, and is susceptible to mistakes. So, an automatic method is required for NT measurement. Materials and Methods: This study proposes a Markov random Field-based approach for contextually segmenting the NT area from foetal pictures and offering a quick and inexpensive diagnostic even during the early stages of pregnancy. Results: Proposed method gives maximum NT thickness error 0.03 and minimum NT thickness error 0.04. Conclusion: The proposed research work developed a prototype for an automated NT thickness measuring system. This study proposes an MRF-based model for segmentation and detection of NT area from foetal pictures which gives error is less than other methods.

Keywords

Nuchal translucency, ultrasound, Markov Random Field, trimester, syndrome

Received 17 January 2024; accepted 24 January 2024

Introduction

Nuchal translucency (NT), as shown in Figure 1, is a liquid fill under the foetus's neck skin that shows sonographically like an anechogenic area (i.e., a black zone in greyscale images) among two echogenic areas (i.e., bright zones). The thickness of the NT, also known as the diameter, is associated not only with Down's Syndrome (DS), but also with other genetic conditions such as Edwards' (trisomy 18), Palau's (trisomy 13), Turner's syndromes, and heart problems.^{1,2}

For almost two decades, researchers have been captivated by the link between increasing foetal neck thickness and genetic and cardiac problems.3 Benacerraf et al.4 reported a study in 1987 that linked DS to higher nuchal fold thickness evaluated during 16 weeks of gestation in 5500 foetuses. Following that, Nicolaides et al.5 reported an increased incidence of chromosomal abnormalities in foetuses with high nuchal fluid buildup at 10-14 weeks gestation. NT is a subcutaneous accumulation of fluid in the foetal neck that is detected by ultrasonography during the initial trimester of pregnancy.6 NT thickness rises with foetal crown-rump length in normal foetuses (CRL). Contributing factors for expanded NT usually involve cardiac dysfunction associated with heart

and great artery abnormalities, venous congestion inside the head and neck, altered extracellular matrix composition, collapse of lymphatic drainage due to atypical or delayed lymphatic system development, or impaired foetal movements, foetal anaemia or hypoproteinaemia, and congenital infection. Table 1 summarises the relationship between NT thickness and the frequency of chromosomal problems, miscarriage, foetal mortality, and significant foetal abnormalities. With increasing NT thickness, the frequency of chromosomal abnormalities grows dramatically. It became obvious very quickly that an elevated NT is not just a diagnostic for chromosomal abnormalities, but also a generic evidence of a disruption in regular early development.

Department of Electronics Engineering, Bharati Vidyapeeth (Deemed to be University) College of Engineering, Pune, Maharashtra, India ²Department of Electronics & Telecommunication, Bharati Vidyapeeth (Deemed to be University) College of Engineering, Pune, Maharashtra, India

Corresponding author:

Kalyani Chaudhari, Department of Electronics Engineering, Bharati Vidyapeeth (Deemed to be University), College of Engineering, Pune, Maharashtra 411043, India. E-mail: kalyanichaudhari1979@gmail.com

 \odot \odot

Creative Commons Non Commercial CC BY-NC: This article is distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 License (http://www.creativecommons.org/licenses/by-nc/4.0/) which permits non-Commercial use, reproduction and distribution of the work without further permission provided the original work is attributed as specified on the SAGE and Open Access pages (https:// us.sagepub.com/en-us/nam/open-access-at-sage).

Accurate measurements are critical because differences of even just a few tenths of a millimetre might have therapeutic implications owing to the influence on the detection rate.7 The positioning of the callipers on the monitor is an important factor in NT measurement quality. Callipers could be adjusted by a small distance of 0.1 mm using current technology. The method for measuring NT, the necessity for proper sonographer training and external quality verification, and the use of NT in successful monitoring for genetic abnormalities are all well established.8 Traditional segmentation algorithms confront several obstacles, including fuzzy edges, intensity uniformity, increased time consumption, and a significant likelihood of inaccuracy in recovering the NT area. Rapid identification of DS would overcome these challenges and give a more accurate and timely diagnosis of the foetus. The objective of this article is to introduce an effective tool to aid in early diagnosis through the automatic quantification of NT; the suggested scheme can identify NT and a way of measuring its thickness without the need for user intervention, thereby eliminating inter- and intraobserver variability.

The following is how the article is structured. The second section provides a summary of the background literature. The third section presents a problem formulation for estimating the NT area using Fuzzy Clustering and MRF. The



Figure I. Nuchal Translucency.¹

segmentation method used to detect the NT edges properly is described. The fourth section presents the experimental outcomes. The fifth section concludes the article.

Literature Survey

There are various publications on the measurement of NT in the literature; Sciortino et al.9 present a novel approach to assist the clinician in both automatically identifying the nuchal area and obtaining an accurate thickness estimate of the NT. Thomas et al.¹⁰ describe a SegNet model based on a Convolutional Neural Network (CNN) and a VGG-16 for semantically separating the NT area from US foetal images and delivering a rapid and economical assessment during the beginning phases of gestation. An AlexNet-based transfer learning technique is used to train the NT segmented areas for the detection of the midsagittal plane, Chudhary et al.11 suggested an automated NT identification approach based on SIFT feature point and GRNN. This non-invasive method is critical not just for NT testing, but also for detecting severe abnormalities and identifying high-risk pregnancies.¹² Ping et al.¹³ investigate the use of ultrasonic assessment of foetal NT width in early gestation and its association with unfavourable pregnancy outcomes. Lee et al. proposed^{14,15} a semi-automatic method of foetal NT thickness measurement that utilises a coherence-enhancing diffusion filtration to improve the frontier and eliminate, accompanied by identification of the NT using optimisation techniques to minimise a cost function that combines intensity, edge strength, and continuity. Catanzariti et al.¹⁶ offer a method that finds the boundary of an NT automatically once a targeted area has been explicitly selected. The approach is based on cost function minimisation and is optimised using the dynamic programming paradigm.

Deng et al.¹⁷ suggested a hierarchical structural model for identifying the NT area automatically. To represent the NT, head, and body of foetuses, three discriminative classifications are initially trained using Gaussian pyramids. The spatial constraints among them are then denoted by a spatial model. Eventually, the suggested model is inferred using dynamic programming and the generalised distance transform, ensuring that the best solution for NT detection is reached. Nirmala et al. reported¹⁸ the NT Width measurement in screening

Table 1. IN I Thickness and the Frequency of Chromosomal Problem	nickness and the Frequency of Chromosomal Pro	oblems ^{2,1}
---	---	-----------------------

	Nuchal Translucency ¹⁰	Major Foetal Abnormalities ¹⁰	Foetal Death ¹⁰	Chromosomal Defects ² [AI_2]					
Ι	>6.5 mm	46.2%	19.0%	64.5%					
2	5.5–6.4 mm	24.2%	10.1%	50.5%					
3	4.5–5.4 mm	18.5%	3.4%	33.3%					
4	3.5–4.4 mm	10.0%	2.7%	21.1%					
5	95th–99th centiles	2.5%	1.3%	3.7%					
6	<95th centile	1.6%	1.3%	0.2%					

first-trimester foetus for DS. The NT zone was segmented using mean shift analysis as well as canny operators, and the precise thickness was calculated using Blob analysis. Liu et al.¹⁹ create a CNN comprising fully linked layers to directly identify the NT area. Explicit identification of other body parts, on the other hand, necessitates extra annotation, development, and processing expenditures. In circumstances of ambiguous head placement or non-standard head-NT relations, it may also create cascading mistakes. In addition, we use U-Net with a tailored design and loss algorithm to provide exact NT segmentation. Finally, principal component analysis is used to compute NT thickness.

Methodology

The work that this article discusses is part of a larger initiative aimed at designing and developing software tools that could simplify the entire procedure of non-invasive screening for the operator. An automated system has several advantages. First, the evaluation would become consistent and repeatable, increasing the overall reliability of the test and perhaps minimising the frequency of intrusive screenings.²⁰ Second, with automated measurement, the time required for a single assessment would be reduced, shortening the time patients must wait before being evaluated while enhancing the number of screens that may be conducted daily.²¹

The goal of image segmentation is to divide the image into a set of meaningful chunks, or regions, based on the intensity values indicated by their pixels. Specifically, segmentation divides the image into a set of *c* crisp regions $\{\mathcal{R}_i\}$ that are maximum linked and each \mathcal{R}_i is consistent based on a set of criteria. Normally, determining whether a pixel belongs to an area is challenging. To cope with such a circumstance, the segmentation procedure employs fuzzy set ideas. FCM (Fuzzy C-mean) (Pal and (Ed.). 1992) is one of the most used fuzzy clustering techniques for medical image segmentation. FCM divides $\{x_k\}_{k=1}^{N}$ into clusters by minimising the objective function provided as:

$$\mathcal{J} = \sum_{i=1}^{c} \sum_{k=1}^{N} u_{ik}^{m} || x_{k} - v_{i} ||^{2}$$
[1]

 x_k = Gray Value associated with k^{th} pixel

 $v_i = i^{th}$ Cluster Center

 u_{ik} = Fuzzy menmbership Value of pixel k

m = Fuzziness Exponent with valuelarger than 1

Mobility of the contour is constrained by the provided image in active contour approaches. If the item being detected is present in the image, the contour will begin above it and move inward along the object's normal until it hits the edge. The LMS, which was first proposed by Osher and Sethian in 1988, employs a zero-level function supplied at time t as a 2D Lipschitz function indicated by $\varphi(x, y)$ representing contour

$$C = \left\{ \frac{(x, y)}{\varphi(x, y)} = 0 \right\}$$
 which evolves through zero-level function

given at time t as $\varphi(x, y)$. The evolution equation for a levelset function may be expressed as a partial differential equation, as follows:

$$\frac{\partial \varphi}{\partial t} = \left| \nabla \varphi \right| \mathcal{F}, \varphi \left(0, x, y \right) = \varphi_0 \left(x, y \right)$$
[2]

In this case, the coordinates $\varphi(0, x, y)$ reflect the starting shape, while \mathcal{F} stands for both external and internal forces.

Proposed Segmentation

Initially, the suggested technique would segment the region of interest (ROI) using a level-set method and fuzzy clustering. In the suggested method, the level-set function is initialised to an initial value using fuzzy clustering. The FCM creates misleading blobs and outliers in the picture. Gaussian filtering is used to isolate these ancillary effects. Following these steps, the results of the fuzzy clustering are used to initialise the level-set function.

$$\mathcal{M}_{i,j}$$
 = Membership Function

 $\mathcal{Z}_{i,j} = \text{ROI} \text{ extracted fromImage}$

A ROI is computed using,

$$\mathcal{Z}_{i,j} = \begin{cases} 1, & u_{ij} > b_0 \\ 0, & otherwise \end{cases}$$

$b_0 \in (0,1)$ is adjustable threshold

The level-set function is set up initially as:

$$\varphi_0(x, y) = 2\varepsilon \left(2\mathcal{Z}_{i, j} - 1\right)$$
[3]

ε = constant meant for regularizing the Dirac function

In traditional level-set approaches, controlling parameters are inputted manually, with values shifting according to the needed application, in order to influence the development of the level-set function. The suggested technique utilises the outputs of fuzzy segmentation to perform an adaptive evaluation of all the necessary controlling parameters. The Heaviside function and the Dirac function are used to assess the area and the length of the contour generated by fuzzy clustering.

$$C_{Area}\left(\varphi \ge 0\right) = \int_{-\Omega}^{\Omega} \mathcal{H}\left(\varphi\left(x, y\right)\right) dx \, dy \qquad [4]$$

$$\mathcal{C}_{Length}\left(\varphi=0\right) = \int_{-\Omega}^{\Omega} \delta_{0}\left(\varphi\left(x,y\right)\right) \left|\nabla\varphi\left(x,y\right)\right| dx \, dy \qquad [5]$$

Here, the Heaviside function may be described as:

$$\mathcal{H}(\varphi) = \begin{cases} 1, & \text{if } \varphi \ge 0\\ 0, & \text{if } \varphi < 0 \end{cases}$$
[6]

and the Dirac function as:

$$\delta_0(\varphi) = \frac{d}{d\varphi} \mathcal{H}(\varphi)$$
 [7]

An increase in balloon pressure, the extraction zone is drawn in by the level-set function, which is generated by the parameter m. Based on the sign of m, the balloon force either contracts or expands the level-set function, eliminating the need to initialise the level-set function at the limits of the needed object. The pressure is represented by the size of m, which is shown to be sufficiently big to squeeze through artificial boundaries. This means that in the suggested technique, the value of m is calculated based on the output of the fuzzy segmentation as:

$$v = 0.5 - u_{i,i}$$
 [8]

$u_{i,j}$ = membership function associated with fuzzy segmentation for every pixel

The Markov Random Field (MRF) is a sophisticated stochastic modelling technique that may be used to efficiently depict the local interactions between the properties of neighbouring pixels. More than that, it may use Bayesian probability distributions to represent the spatial relationship between neighbouring pixels. Segmentation is performed using the maximum posterior probability (MAP) of the labelling space provided by the image data.²² Energy maximisation problems are at the heart of the MRF MAP specifications. In addition to the combinatorial nature of the maximisation problem, the nonconvexity of the energy function and the presence of several local minima in the image's solution space make it difficult to achieve a global maximum. The innovative Hidden Markov Random Field (HMRF) method was developed as a spinoff of the Hidden Markov Model,²³ a Markov chain-based algorithm. As a graphical probability prototype, HMRF does not directly discover the proper states but rather estimates them via a network of observations.

An observable random field is present in HMRF models,

$$a = (a_1, \ldots, a_N)$$

a_i = Feature Value of Pixel

Which seeks to deduce an underlying stochastic field

Hidden Random Field =
$$(b_i, \ldots, b_N)$$

$$b_i \in \mathcal{K}$$

$$\mathcal{K} =$$
set of all possible labels.

 $b_i = \text{configuration of labels}$

If the following Markovian condition is true, then the label field a_i with regard to its neighbourhood system is an MRF.

$$\rho(\mathbf{a}|\mathbf{b}) = \prod_{i \in S} \rho(a_i|\mathbf{b}_i)$$
[9]

S Spot related to another one another with neighbourhood System

$$\mathcal{N} = \{\mathcal{N}_i, i \in \mathcal{S}\}$$

$$\mathcal{N}_i = \text{pots adjacent to pixel } i$$

b_{N_i} = Neighbourhood Configuration

MRF's Equation z_1 for conditional independence looks like this

$$b(a \mid b) = \rho(a \mid b)\rho(x) = \rho(b) \prod_{i \in S} \rho(a_i \mid b_i)$$
[10]

$$\rho(a_i, b_i \mid b_{Ni}) = \rho(a_i \mid b_i) \rho(b_i \mid b_{Ni})$$
[11]

Assuming *a* is selected at random from a probability distribution $f(a;k,\theta)$, for $k \in \mathcal{K}$, the marginal probability of a_i is given by:

$$\rho(a_i \mid b_{Ni}, \theta) = \sum_{k \in \mathcal{K}} \rho(a_i, k \mid b_{Ni}, \theta) = f(a_i, \theta_k) \rho(k \mid b_{Ni})$$
[12]

Measurement of the HMRF using the EM algorithm:

$$\hat{b} = \arg\max\left\{\rho(a \mid b, \Theta)\rho(b)\right\}$$
[13]

A Gibbs distribution may be thought of as an alternative way to describe an MRF:

$$\rho(b) = z^{-1} exp\left(-\mathcal{V}_{(b)}\right) \qquad [14]$$

z: normalization constant

$$\mathcal{V}_{(b)}$$
 = potential function

In Equation [12], the prior probability $\rho(b)$ is a Gibbs distribution, and its joint probability is expressed as

$$\rho(a \mid b, \Theta) = \prod_{i} \rho(a_i \mid b, \Theta) = \prod_{i} \rho(a_i \mid b_i, \theta_{bi}) \qquad [15]$$

 $\rho(a_i | \mathbf{b}_i, \theta_{bi})$: Gaussian distribution with parameters

$$\theta_{bi} = \left(\mu_{bi}, \Sigma_{bi}\right)$$

To summarise the HMRF-EM method in a nutshell:

Step 1: Initialise parameter set.

Step 2: The second stage entails determining the probability distribution. $\rho(a_i|b_i, \theta_{bi})$.

Step 3: Using $\Theta^{(t)}$ to calculate labels by MAP estimation:

$$b^{(t)} = \arg \max_{b \in \mathcal{B}} \left\{ \rho\left(a \mid b, \Theta^{(t)}\right) \rho(b) \right\}$$
$$= \arg \min_{b \in \mathcal{B}} \left\{ v\left(a \mid b, \Theta^{(t)}\right) + v(b) \right\}$$
[16]

Step 4: The posterior probability distribution is calculated $k \in \mathcal{K}$ and pixels a_i ,

$$\rho^{(t)}\left(k \mid a_{i}\right) = \frac{\mathcal{N}\left(a_{i}, \theta_{i}\right)\rho\left(k \mid b_{N_{i}}^{(t)}\right)}{\rho^{(t)}b_{i}}$$
[17]

 $\mathcal{N}(a_i, \theta_i)$ = Gaussian Distribution of a_i with

$$\theta_{k} = \left(\mu_{k}, \Sigma_{k}\right)$$

$$\rho\left(k \mid b_{\mathcal{N}_{i}}^{(t)}\right) = \frac{1}{z} \exp\left(-\sum_{j \in \mathcal{N}_{i}} \mathcal{V}_{c}\left(k, b_{j}^{(t)}\right)\right)$$

Step 5: Parameters of the Gaussian distribution are modified by $\rho(k | b_{N_i}^{(t)})$.

An initial segmentation is created in the HMRF method for image segmentation by applying a Level set based on the grey-level values of pixels. Significantly fewer iterations are required to produce final labels using this method compared to the MRF-MAP method, demonstrating the importance of this procedure.

The suggested technique uses the energy functions created for the integration of Clustering based Level-set and MRF to establish the first segmentation of pictures. The standard Potts model presents the regional prior energy function, often known as the potential function, as,

$$\varepsilon = -\sum_{j \in \mathcal{N}_i} \mathcal{V}_c\left(k, b_j^{(t)}\right)$$
[18]

 $\mathcal{V}_c =$ Kronecker delta which equals 1 for $k = b_j^{(t)}$ and being 0 otherwise.

As an added complication, the posterior energy function requires modelling the image's probability density function.

NT thickness measurement

In the first three months of pregnancy, the NT thickness is a key diagnostic indicator. While NT thickness is relatively insensitive, this is not the case.¹⁸ Fluid accumulates in the nuchal region of the foetus, hence the NT size is much more susceptible to the deposition than that of the NT thickness. The suggested method may also be used to calculate the size of the NT. To successfully extract the NT contour from the ultrasound picture, the suggested method is used. Because of this, calculating the parameters is straightforward and achievable. Before we can measure the NT thickness, we need to locate its limits. We offer the following approach to calculating the NT's margins inside the chosen ROI. Take into account a ROI with N × M dimensions. The NT region is measured by the number of pixels that fall inside the NT boundary. Edge detection is used on this ROI to identify the ceiling and floor. The objective of the thickness evaluation is to provide a good estimate for the largest possible separation of the two boundaries identified.

Experiment

The proposed research work data collection includes 200 ultrasound images from the first trimester of pregnancy. Dataset is collected from women of the age group 25–40. A section of the NT was annotated by specialists. The discovered NT edges are shown by the yellow marker in (Figure 2).

The error between Measured and Automatic NT thickness measurements (Maximum) is shown in Table 2.

Figure 3 shows that the suggested approach has a smaller margin of error relative to manual measurement than the other methods.

The error between Measured and Automatic NT thickness measurements (Minimum) is shown in Table 3.



Figure 2. Input Image and NT Detected Image.



Figure 3. Comparison of Maximum NT Thickness.

Method	Measured Maximum NT Thickness in mm	Automatic Maximum NT Thickness in mm	Error in mm	Method	Measured Maximum NT Thickness in mm	Automatic Maximum NT Thickness in mm	Error in mm	
Proposed	2.93	2.9	0.03	Proposed	0.75	0.7	0.04	
SegNet ¹⁹	2.93	2.7	0.23	SegNet ¹⁹	0.75	0.5	0.2	
GRNN	2.93	2.84	0.09	GRNN ¹¹	0.75	0.27	0.47	
RBED ¹⁸	2.93	2.7	0.23	RBED ¹⁸	0.75	0.49	0.25	
HSM ¹⁷	2.93	2.48	0.45	HSM ¹⁷	0.75	0.52	0.22	

 Table 2. Error Between Measured and Automatic NT Thickness

 Measurements (Maximum), 11,17–19

 Table 3. Error Between Measured and Automatic NT Thickness

 Measurements (Minimum).^{11–17–19}



Figure 4. Comparison of Minimum NT Thickness.

The minimal thickness of segmented NT is shown in Figure 4. The above Figure 4 shows that the suggested approach has a smaller margin of error relative to manual measurement than the other methods.

Conclusion

Transabdominal or transvaginal ultrasonography is used to determine NT thickness in the sagittal region of foetuses. Nowadays, clinicians are typically in charge of measuring the NT thickness. This necessitates highly competent operators and results in data unpredictability. There has only been a small investigation on an automated computerised assessment of foetal NT. The proposed research work developed a prototype for an automated NT thickness measuring system. This study proposes an MRF-based model for conceptually segmenting the NT area from foetal pictures and offering a quick and inexpensive diagnostic during the early phases of pregnancy.

Future Scope

As the proposed method is computationally complex the automated determination of the appropriate number of states for the above model, or the number of area clusters produced by the image segmentation process, is one of their main limitations. So in the future researcher can upgrade this using deep learning methods. As well as foetal picture size and angle departure from the mid-sagittal plane have an impact on the NT measures; so the researcher has to take care of it.

Acknowledgments

This work would not have been possible without the contributions of other researchers in the area. I am grateful to each and every one of them, especially to Apex Diagnostic Centre, Pune, for supplying a database and valuable advice, and to the gynaecologist, obstetrician, and infertility expert Dr Sonali Chaudhari for her support.

Declaration of Conflicting Interests

The authors declared no potential conflicts of interest with respect to the research, authorship and/or publication of this article.

Funding

The authors received no financial support for the research, authorship and/or publication of this article.

References

- 1. Woodward PJ, Kennedy A, Roya S. Diagnostic imaging: obstetrics. *Diagnostic Imaging Book*. 3rd ed. 2017.
- 2. FMF, Fetal Medicine Foundation nuchal translucency. http:// www.fetalmedicine.org/

- Nicolaides KH, Azar G, Byrne D, et al. Fetal nuchal translucency: ultrasound screening for chromosomal defects in first trimester of pregnancy. *BMJ* 1992; 304(6831): 867–869. https:// doi.org/10.1136/bmj.304.6831.867. PMID 1392745.
- Benacerraf BR, Gelman R, Frigoletto FD. Sonographic identification of second-trimester fetuses with Down's syndrome. N Engl J Med 1987; 317(22): 1371–1376. https://doi.org/10.1056/ NEJM198711263172203. PMID 2960895.
- Nicolaides KH, Azar G, Byrne D, et al. Fetal nuchal translucency: ultrasound screening for chromosomal defects in first trimester of pregnancy. *BMJ* 1992; 304(6831): 867–869. https:// doi.org/10.1136/bmj.304.6831.867. PMID 1392745.
- Souka AP, Von Kaisenberg CS, Hyett JA, et al. Increased nuchal translucency with normal karyotype. *Am J Obstet Gynecol* 2005; 192(4): 1005–1021. https://doi.org/10.1016/j. ajog.2004.12.093. PMID 15846173.
- Kagan KO, Wright D, Etchegaray A, et al. Effect of deviation of nuchal translucency measurements on the performance of screening for trisomy 21. *Ultrasound Obstet Gynecol* 2009; 33: 657–664.
- Nicolaides KH. Nuchal translucency and other first-trimester sonographic markers of chromosomal abnormalities. *Am J Obstet Gynecol* 2004; 191(1): 45–67. https://doi.org/10.1016/j. ajog.2004.03.090. PMID 15295343.
- Sciortino G, Tegolo D, Valenti C. Automatic detection and measurement of nuchal translucency. *Comput Biol Med* 2017; 82: 12–20. https://doi.org/10.1016/j.compbiomed.2017.01.008. PMID 28126630.
- Thomas MC, Arjunan SP. Deep learning measurement model to segment the nuchal translucency region for the early identification of Down syndrome. *Meas Sci Rev* 2022; 22(4): 187–192. https://doi.org/10.2478/msr-2022-0023
- Chudhary K, Oza S. Ultrasound image based fully-automated nuchal translucency segmentation and thickness measurement. *Int J Nonlinear Anal Appl* 2021; 12: 1573–1583. https://doi. org/10.22075/ijnaa.2021.5830
- Mulani AO, Mane PB Watermarking and cryptography based image authentication on reconfigurable platform. *Bulletin EEI* 2017; 6(2): 181–187. https://doi.org/10.11591/eei.v6i2.651
- Ping FU, Wencui WU. Value of ultrasonic measurement of fetal nuchal translucency thickness in early pregnancy and its

relationship with adverse pregnancy outcomes. *J Clin Med Pract* 2021; 25(15): 93–97. https://doi.org/10.7619/jcmp.20211030

- Lee YB, Kim MJ, Kim MH. Robust border enhancement and detection for measurement of fetal nuchal translucency in ultrasound images. *Med Biol Eng Comput* 2007; 45(11): 1143–1152. https://doi.org/10.1007/s11517-007-0225-7. PMID 17657519
- Mulani AO, Birajadar G, Ivković N, et al. Deep learning based detection of dermatological diseases using convolutional neural networks and decision trees. *Trait Signal* 2023; 40(6): 2819– 2825. https://doi.org/10.18280/ts.400642
- Catanzariti E, Fusco G, Isgrò F, et al. A semi-automated method for the measurement of the fetal nuchal translucency in ultrasound images. In:ICIAP 2009. Lecture notes in computer science (ed Foggia P, Sansone C, Vento M), 2009, Vol. 5716, pp. 613–622, Berlin, Heidelberg: Springer. https://doi. org/10.1007/978-3-642-04146-4_66
- Deng Y, Wang Y, Chen P. Automated detection of fetal nuchal translucency based on hierarchical structural model. In: 23rd International Symposium on Computer-Based Medical Systems (CBMS), 2010, Vol. 2010, pp. 78–84. IEEE Publications. https://doi.org/10.1109/CBMS.2010.6042618
- Nirmala S, Palanisamy V. Measurement of nuchal translucency thickness for detection of chromosomal abnormalities using first trimester ultrasound fetal images. *Int J Comput Sci Inf Sec* 2009; 6(3).
- Liu T, Xu M, Zhang Z, et al. Direct detection and measurement of nuchal translucency with neural networks from ultrasound images. *Lect Notes Comput Sci* 2019; 11798; 20–28. https://doi. org/1007/978-3-030-32875-7 3.
- Shinde R, Mulani AO. Analysis of biomedical image using wavelet transform. *Int J Recent Innov Trends Technol* 2015; 2(7).
- Mane DP, Mulani AO. High throughput and area efficient FPGA implementation of AES algorithm. *Int J Eng Adv Technol* 2019; 8(4).
- Montenegro N, Matias A, Areias JC, et al. Increased fetal nuchal translucency: possible involvement of early cardiac failure. *Ultrasound Obstet Gynecol* 1997; 10(4): 265–268. https://doi. org/10.1046/j.1469-0705.1997.10040265.x. PMID 9383878.
- Zhang T, Xia Y, Feng DD. An evolutionary HMRF approach to brain MR image segmentation using clonal selection algorithm. *IFAC Proc Vol* 2012; 45(18): 6–11. https://doi. org/10.3182/20120829-3-HU-2029.00092