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Published in:
Proceedings of Ultrasonics Symposium

Link to article, DOI:
[10.1109/ULTSYM.1990.171635](https://doi.org/10.1109/ULTSYM.1990.171635)

Publication date:
1990

Document Version
Publisher's PDF, also known as Version of record

[Link back to DTU Orbit](#)

Citation (APA):
Jensen, J. A. (1990). Deconvolution of in vivo ultrasound images. In *Proceedings of Ultrasonics Symposium* (Vol. Volume 3, pp. 1581-1587). IEEE. <https://doi.org/10.1109/ULTSYM.1990.171635>

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DECONVOLUTION OF *IN VIVO* ULTRASOUND IMAGES

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Abstract

The appearance of an ultrasound image is intimately linked to the nature of the emitted pressure field and its distortion down through the tissue. The pulse field is temporally and spatially varying due to focusing and the dispersive attenuation.

The influence of the pulse and attenuation should be removed from the picture in order to display a more consistent and uniform image.

This paper describes an algorithm to remove the influence of the attenuated pulse on the image. The algorithm takes into account the varying pulse, noise in the acquired signal and the changing reflectivity in the tissue. Both one- and two-dimensional processing can be implemented. The algorithm relies on prior knowledge of the pulse, and of the covariance of the noise and the reflections. Algorithms to estimate these are given.

Examples of use are presented. One-dimensionally processed images of the kidney and a foetus are shown.

1 Introduction

Ultrasound is routinely used in hospitals for the investigation of nearly all soft tissue structures of the human body. Advanced transducers (phased and linear array) are used in order to acquire sharp images. Improved resolution is steadily sought by increasing the center frequency and bandwidth of the emitted pulse. Increasing the center frequency, however, limits the depth of penetration due to the dispersive attenuation of the pulse in the tissue. This limits the resolution attainable, and to further increase resolution, or depth of penetration for the same resolution, we must turn to digital processing, *i.e.* deconvolution.

Several authors have investigated techniques for enhancing the resolution of ultrasound images. Fatemi and Kak [1] derived an expression for the received pressure field and suggested use of a Wiener filter [2]. This has also been suggested in [3], [4]. Slightly different, but conceptually similar methods have been suggested in [5], [6],

[7] [8]. These estimators use, however, fixed parameters for the emitted pulse and can not take into account the spatially varying source wavelet and the dispersive attenuation, which are important factors for *in vivo* images. We therefore turn to other, more advanced estimators.

In this paper we will devise an algorithm, based on physical reasoning, that can take into account variations in pulse shape, signal-to-noise ratio and reflection strength. It is based on Mendel's fixed-interval deconvolution algorithm [9], [10], which calculates an optimal, minimum variance estimate of the reflections. As this estimator can handle varying parameters optimally, the original problem is transformed into finding parameters to describe the pulse and covariances for the noise and the reflections. These parameter estimators should take into account temporal and spatial variations, and section 4 and 5 will explain how to do this.

Examples of the use of the combined algorithm are shown both for data acquired from a tissue mimicking phantom and for clinical *in vivo* data. These demonstrate the algorithm's capability to increase the resolution and to handle variations in reflection strength. The effect of non-linearities emanating from the data acquisition and ways of avoiding them are discussed.

2 The general algorithm

It can be shown that the received pressure field can be calculated by [11]:

$$p_r(\vec{r}_2, t) = v_{pe}(t) \star_t f_m(\vec{r}_1) \star_r h_{pe}(\vec{r}_1, \vec{r}_2, t) \quad (1)$$

where $\star_t$ denotes a temporal and $\star_r$ a spatial convolution. v_{pe} is the pulse-echo wavelet, that accounts for the transducer excitation and the electro-mechanical impulse response during emission and reception of the pulse. f_m accounts for inhomogeneities in the tissue due to density and propagation velocity perturbations, which gives rise to the scattered signal. h_{pe} is the modified pulse-echo spatial impulse response, that relates the

transducer geometry to the spatial extent of the scattered field. Explicitly written out it is:

$$h_{pe}(\vec{r}_1, \vec{r}_2, t) = \frac{1}{c_0^2} \frac{\partial h(\vec{r}_1, \vec{r}_2, t)}{\partial t} \star_t \frac{\partial h(\vec{r}_1, \vec{r}_2, t)}{\partial t} \quad (2)$$

$\vec{r}_1$ denotes the position of the scatterer and $\vec{r}_2$ the position of the transducer. h is the spatial impulse response for the transducer geometry as calculated by the Tupholme-Stepanishen method [12],[13],[14]. Equation (1) was derived under the assumption of weak scattering by density and propagation velocity perturbations in the tissue, and absorption and multiple scattering were neglected. The dispersive attenuation can be lumped into v_{pe} , that then is substituted by a pulse, $V_{pea}(t, \vec{r}_1, \vec{r}_2)$ that depends on distance.

Expression (1) consists of three distinct terms. The signal of interest, and the one that should be displayed in medical ultrasound, is $f_m(\vec{r}_1)$. We, however, measure a time and spatially smoothed version of this, which obscures the finer details in the image. The smoothing consists of a convolution in time with a wavelet $v_{pea}(t, \vec{r}_1, \vec{r}_2)$, and a spatial convolution with a spatially varying $h_{pe}(\vec{r}_1, \vec{r}_2, t)$.

It is necessary to employ digital signal processing to make a better estimation of f_m . Thus, (1) is reformulated in a discrete form.

$$y(\vec{r}_{d_2}, k) = v_{pea}(k, \vec{r}_{d_1}, \vec{r}_{d_2}) \star_k f_m(\vec{r}_{d_1}) \star_{\vec{r}_d} h_{pe}(\vec{r}_{d_1}, \vec{r}_{d_2}, k) + n(\vec{r}_{d_2}, k) \quad (3)$$

k is the discrete time variable, and $\vec{r}_d$ is a discrete vector. A noise term $n(\vec{r}_{d_2}, t)$ has been added to explain the inevitable noise in the measured signal. The term accounts for both electrical noise from amplifiers and for physical effects not explained by the convolution model.

We now seek a procedure to obtain an estimate of $f_m(\vec{r}_{d_1})$. This can formally be stated as [15]:

$$\hat{f}(\vec{r}_{d_1}) = \mathcal{F} [y(\vec{r}_{d_2}, k), v_{pea}(k, \vec{r}_{d_1}, \vec{r}_{d_2}), h_{pe}(\vec{r}_{d_1}, \vec{r}_{d_2}, k), n(\vec{r}_{d_2}, k)] \quad (4)$$

The estimate $f_m(\vec{r}_{d_1})$ is a function of the measured signal $y(\vec{r}_{d_2}, k)$ and some knowledge, measured or estimated, about v_{pea} , h_{pe} , and n .

The function $\mathcal{F}$ can be a linear or non-linear mapping from y , v_{pea} , h_{pe} , and n to $\hat{f}_m$. It can be evaluated directly or through iterative techniques. One of the advantages of ultrasound B-mode systems is real-time image formation. This precludes an iterative search scheme, as in a real-time system only a fixed amount of calculations can be carried out per sample. Rather the estimate

$\hat{f}_m(\vec{r}_{d_1})$ is based on some estimate of the internal state of the system so that

$$\begin{aligned} \hat{S}(\vec{r}_{d_2}, k) &= \mathcal{H} [y(\vec{r}_{d_2}, k), v_{pea}(k, \vec{r}_{d_1}, \vec{r}_{d_2}), \\ &\quad h_{pe}(\vec{r}_{d_1}, \vec{r}_{d_2}, k), n(\vec{r}_{d_2}, k)] \\ \hat{f}_m(\vec{r}_{d_1}) &= \mathcal{G} [\hat{S}(\vec{r}_{d_2}, k)] \end{aligned} \quad (5)$$

where $\mathcal{H}$ linearly relates y , v_{pea} , h_{pe} , and n to $\hat{S}$, and $\mathcal{G}$ linearly relates $\hat{S}$ to $\hat{f}_m$. S is a vector of fixed dimension that represents the state of the system. $\mathcal{H}$ and $\mathcal{G}$ can be evaluated with a fixed and a priori known number of calculations. This allows for a time-varying model, where each sample can be treated in a fixed, predetermined interval of time [15].

3 The deconvolution algorithm

Eq. (5) explains in general terms how to remove v_{pea} and h_{pe} from y to obtain f_m . We will now detail how to perform this deconvolution, when the parameters are known.

The algorithm used is Mendel's fixed-interval deconvolution estimator. It consists of two parts: first a Kalman filtering is performed on the data time-recursively, and then a subsequent estimation step performed backwards time-recursively. The result is a minimum variance, fixed-interval estimate of the reflection signal. The algorithm can be used on one A-line or on a number of A-lines at a time, and thus perform two-dimensional deconvolution.

For easy reference the basic equations are now given. The Kalman filter is based on the following time-varying state-space model:

$$\begin{aligned} x(k+1) &= \Phi(k+1, k)x(k) + \Gamma(k+1, k)w(k+1) \\ z(k) &= H^T(k)x(k) + n(k) \end{aligned} \quad (6)$$

$z(k)$ is the measured signal, $w(k+1)$ is the "tissue" reflection signal, $x(k)$ the state vector, and $n(k)$ the measurement noise. Φ is the state transition matrix, Γ the input distribution matrix, and H the output matrix. Note the direct coupling between the input $w(k)$ and $z(k)$. The equations describe a time-varying model, for which a new set of coefficients in the state matrices and vectors can be used for each time instance. It is assumed that the $w(k+1)$ and $n(k)$ are zero mean, Gaussian signals with a covariance of $Q(k+1)$ and $R_n(k)$.

The Kalman filter is, stated in the predictor-corrector form:

Predictor equations

$$\hat{x}(k+1 | k) = \Phi(k+1, k)\hat{x}(k | k) \quad (7)$$

$$P(k+1|k) = \Phi(k+1, k)P(k|k)\Phi^T(k+1, k) + \Gamma(k+1, k)Q(k+1)\Gamma^T(k+1, k) \quad \varphi^T(k) = [-y(k-1), \dots, -y(k-n_a), e(k-1), \dots, e(k-n_c)] \quad (12)$$

Then the output is written as:

Corrector equations

$$y(k) = \varphi^T(k)\theta + e(k) \quad (13)$$

$$\begin{aligned} K(k+1) &= P(k+1|k)H(k+1) \cdot \\ &\quad [H^T(k+1)P(k+1|k)H(k+1) \\ &\quad + R_n(k+1)]^{-1} \\ \hat{z}(k+1|k) &= z(k+1) - H^T(k+1)\hat{x}(k+1|k) \\ \hat{x}(k+1|k+1) &= \hat{x}(k+1|k) + K(k+1)\hat{z}(k+1|k) \\ P(k+1|k+1) &= [I - K(k+1)H^T(k+1)] \cdot \\ &\quad P(k+1|k) \end{aligned} \quad (8)$$

and the prediction and prediction error as:

$$\begin{aligned} \hat{y}(k|\theta) &= \varphi^T(k)\theta \\ \hat{e}(k, \theta) &= y(k) - \hat{y}(k|\theta) \end{aligned} \quad (14)$$

The parameters are found by minimizing the magnitude of the prediction error. This is formulated as:

$$V_N(\theta, \mathcal{Z}(N)) = \frac{1}{N} \sum_{k=1}^N \hat{e}(k, \theta) \hat{e}(k, \theta) \quad (15)$$

$$\hat{\theta}_N = \underset{\theta \in D_{\mathcal{M}}}{\text{Arg min}} V_N(\theta, \mathcal{Z}(N)) \quad (16)$$

$P(k+1|k)$ is the a-priori state covariance, and $P(k+1|k+1)$ the a-posteriori covariance. $K(k)$ is the Kalman gain.

The backwards time-recursive estimation step can be expressed as:

$$\begin{aligned} \hat{w}(k+1|N) &= Q(k+1)\Gamma^T(k+1, k)r(k+1|N) \\ r(k|N) &= H(k)[H^T(k)P(k|k-1)H(k) \\ &\quad + R_n(k)]^{-1} \hat{z}(k|k-1) + (\Phi(k+1, k) \\ &\quad [I - K(k)H^T(k)])^T r(k+1|N) \end{aligned} \quad (9)$$

$\hat{w}(k+1|N)$ denotes the fixed-interval estimate of $w(k+1)$. $r(k+1|N)$ is the residual state vector [9].

The algorithm can, based on all samples in one or more A-lines, estimate the reflections. The pulse parameters and the covariances can vary from sample to sample and the pulse can be non-minimum phase [9].

4 Fixed-interval pulse estimator

In (5) knowledge of v_{pea} and h_{pe} is needed. v_{pea} must be estimated because, in general, dispersive attenuation is unknown. h_{pe} can be calculated from a priori knowledge of the transducer geometry.

A representative model for v_{pea} is the ARMA (AutoRegressive Moving Average) model given by:

$$(1 + a_1 q^{-1} \dots + a_{n_a} q^{-n_a})y(k) = (1 + c_1 q^{-1} \dots + c_{n_c} q^{-n_c})e(k) \quad (10)$$

where $y(k)$ is the measured signal, $e(k)$ the reflection signal, q^{-1} the unit backward shift operator, and $a_1, \dots, a_{n_a}$ and $c_1, \dots, c_{n_c}$ are the ARMA coefficients.

The parameters to be estimated, and the data values, are, for notational convenience, organised in a parameter vector, θ , and a regression vector, φ .

$$\theta^T = [a_1, a_2, \dots, a_{n_a}, c_1, c_2, \dots, c_{n_c}] \quad (11)$$

stating that V_N is minimized with respect to θ under the constraint that the parameters belong to the set of stable models $D_{\mathcal{M}}$. $\mathcal{Z}(N)$ denotes the set of measurements $\{y(1), y(2), \dots, y(N)\}$. This parameter estimation method is called the prediction error method (PEM) [15], [16].

Using an iterative search algorithm employing the Gauss-Newton search gradient, we arrive at the following equations [16]:

START:

$$\begin{aligned} \psi(k, \theta) &= [1 - C(q)]\varphi(k) \\ V_N^{(i)}(\theta, \mathcal{Z}(N)) &= -\frac{1}{N} \sum_{k=1}^N \psi^T(k, \theta) \hat{e}^{(i)}(k, \theta) \\ V_N^{\prime\prime(i)}(\theta, \mathcal{Z}(N)) &= \frac{1}{N} \sum_{k=1}^N \psi^T(k, \theta) \psi(k, \theta) \\ \alpha &= 1 \end{aligned}$$

ITERATE:

$$\hat{\theta}_N^{(i+1)} = \hat{\theta}_N^{(i)} + \alpha [V_N^{\prime\prime(i)}(\theta, \mathcal{Z}(N))]^{-1} V_N^{\prime(i)}(\theta, \mathcal{Z}(N)) \quad (17)$$

Mirror zeros and poles into the unit circle.

$$\begin{aligned} \hat{e}^{(i+1)}(k, \theta) &= y(k) - \varphi^T(k)\theta_N^{(i+1)} \\ V_N^{(i+1)}(\theta, \mathcal{Z}(N)) &= \frac{1}{N} \sum_{k=1}^N \hat{e}^{(i+1)}(k, \theta) \hat{e}^{(i+1)}(k, \theta) \end{aligned}$$

If $V_N^{(i+1)} < V_N^{(i)}$ then go to START. else $\alpha = \alpha/2$ and go to ITERATE with the new α until tried ten times. If $\|[V_N^{\prime\prime(i)}(\theta, \mathcal{Z}(N))]^{-1} V_N^{\prime(i)}(\theta, \mathcal{Z}(N))\| < 0.01$ or $i = 10$ then stop

i is the iteration step, and $\|\bullet\|$ denotes the norm of the vector.

Some initial parameter and prediction error estimates must be supplied. They are calculated by the least-squares method.

1. Fit an AR-model of order $n_a + n_c$ to the data

$$\text{Model: } A_1(q)y(k) = e(k)$$

$$\hat{\theta}_{\text{LS1}} = \left[\frac{1}{N} \sum_{k=1}^N \varphi_1(k) \varphi_1^T(k) \right]^{-1} \frac{1}{N} \sum_{k=1}^N \varphi_1(k) y(k) \quad (18)$$

2. Calculate the initial estimate of the prediction error

$$\hat{e}(k, \theta_{\text{LS1}}) = y(k) - \varphi_1^T(k) \hat{\theta}_{\text{LS1}} \quad (19)$$

3. Make a least squares fit to an ARX-model of order n_a and n_c , where the external input is $\hat{e}(k, \theta_{\text{LS1}})$

$$\text{Model: } A(q)y(k) = C(q)e(k)$$

This gives the initial parameter estimate for θ and the prediction error.

This estimator gives one set of parameters for a segment of an A-line. It can be reformulated to give one set for each sample [16] tracking the slow variation in the pulse. We, however, prefer to divide the A-line into overlapping segments, and then use parameters from the previous segment to initialize the next segment. This has given more consistent results than using the recursive prediction error method.

5 Covariance estimation

The deconvolution algorithm uses the covariance of the noise and of the reflections. The noise is due to the analog amplifiers, the time gain amplifier and quantization of the signal. It can be modeled as a fixed noise source added to the noise from the TGC amplifier that varies with its gain factor. So the noise covariance can be found from previous measurement of the noise sources and the gain factor actually used for the acquired data,

When the signal-to-noise ratio is good we can approximate the measured signal by the ARMA model in (10). The reflections can then be approximately obtained by performing a predictive deconvolution by the ARMA filter and then finding the covariance. To reduce the variance of the covariance estimate several adjacent lines are used, and each line is weighted by a Hamming window.

Figure 1: Position of wires.

6 Phantom examples

We will now show results from deconvolving signals obtained from a tissue phantom (Nuclear Associates). As we have not been able to create an invertible model for the pulse-echo spatial impulse response, only a 1D deconvolution will be performed with a fixed, estimated wavelet. As the data segment only covers a relatively small depth, a fixed value of Q_{v_r} and R_{v_n} will be used. The values are determined by the method described in the previous section.

The area scanned contained wires as shown in Fig. 1, embedded in a substrate which generates a speckle signal.

The transducer used was the Brüel & Kjær 8526. The focal point is at 5 cm and the nominal frequency is 3.5 MHz. The sampling frequency was 25 MHz.

The ARMA(6,5) wavelet used by the deconvolution algorithm was estimated from one of the acquired lines which contained only a speckle signal. The wavelet is shown in Fig. 2.

Using the approach in section 5, the ratio Q_{v_r}/R_{v_n} was estimated to be 9.2, and this was used in the deconvolution. The signal-to-noise ratio was estimated to 43.

Fig.3 shows the logarithmic envelope of the signals, found by Hilbert transformation. The picture at the top is the normally processed picture, as displayed on a modern scanner.

Fig. 3 shows that the speckle pattern is suppressed and the wires are more distinct than in the traditional picture. This makes it easier to distinguish between speckular scatterers and reflections as the contrast in the picture is enhanced.

Figure 2: Estimated wavelet for the 8526 Brüel & Kjær transducer.

7 *In vivo* examples

We now present two examples with clinical *in vivo* data. The data were acquired by our own prototype sampling system that uses a 12 bits, 20 MHz converter. The measurements were performed at Herlev University Hospital in Denmark with a Brüel & Kjær 1846 scanner with a 8526 3 MHz sector scan transducer.

The first image shown in Fig. 4 is the lower part of the right kidney in a longitudinal scan.

Although the change is not as pronounced as for the phantom data, we do see an improvement. The speckle pattern is finer, and it seems easier to identify the exact boundary of the kidney. The internal structure of the kidney also seems more clearly identified from the speckle noise.

The second example shown in Fig. 5 is a 13 weeks foetus. The head, limbs, and spinal cord can clearly be identified. The deconvolved picture shows the same changes, a slightly finer speckle pattern with, apparently easier identification of the extent of the placenta.

One feature, however, stands out in the deconvolved image; the diamond shaped, white area behind the head. This is due to overload of the input amplifier in the scanner due to a very large reflection. The algorithm relies on a linearity assumption, which is violated here. Thus, the measurement process should ensure that no non-linearities occur. This should not be done by adjusting the TGC for the whole picture, rather an adaptive TGC for each individual line should be used. Thereby a relatively large signal-to-noise ratio can be obtained for the whole image.

Figure 3: Normal and deconvolved response for image measured by the Brüel & Kjær 8526 transducer. Axial distance 30 mm and lateral distance 50 mm. Linear scan.

Figure 4: Normal (left) and deconvolved (right) image of longitudinal scan of right kidney. 1 cm between markers.

Figure 5: Normal (left) and deconvolved (right) image of 13 weeks old foetus. 1 cm between markers.

8 Conclusion

An optimal, fixed-interval, minimum variance algorithm for increasing the resolution of medical ultrasound pictures was given. It can take into account the spatial and temporal change in the emitted and received pressure field and changes in the noise and reflection covariance.

The algorithm uses knowledge of the interrogating pulse and of the covariances, and estimators for the one-dimensional pulse and for the covariances were given.

The algorithm can handle two-dimensional deconvolution, but currently we have not been able to obtain a stable and invertible 2D model for the pressure field.

Examples of 1D deconvoluted pictures of phantom data and *in vivo* data were given. They showed, especially for the phantom data, an increased contrast and resolution. Problems were encountered with *in vivo* images due to non-linearities in the acquired signals. It is suggested that this can be solved by using adaptive time gain compensation selected for each line in the image.

Acknowledgement

The research was partly funded by the Danish Technical Research Council, grant 16-4218.E, Brüel & Kjær A/S, Novo's Foundation, H.C. Ørsted's Foundation and Trane's Foundation.

M.Sc. Jan Mathorne constructed the sampling system and M.Sc. Torben Gravesen and Dr. Bjarne Stage wrote the programs for the data acquisition.

Dr. Sidney Leeman, King's College London is thanked for proofreading the manuscript.

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