



AU9465736

(12) PATENT ABRIDGMENT **(11) Document No. AU-B-65736/94**
(19) AUSTRALIAN PATENT OFFICE **(10) Acceptance No. 674576**

- (54) Title
DECONVOLUTION OF MLS RESPONSE DATA
- International Patent Classification(s)
(51)⁶ **G06F 017/15** **G06F 159/00**
- (21) Application No. : **65736/94** (22) Application Date : **22.04.94**
- (87) PCT Publication Number : **WO94/25925**
- (30) Priority Data
- (31) Number (32) Date (33) Country
9308715 **27.04.93** **GB UNITED KINGDOM**
- (43) Publication Date : **21.11.94**
- (44) Publication Date of Accepted Application : **02.01.97**
- (71) Applicant(s)
MEDICAL RESEARCH COUNCIL
- (72) Inventor(s)
ARTHUR ROGER DAVID THORNTON; JOHN DAVID CHAMBERS; TIMOTHY JOHN FOLKARD
- (74) Attorney or Agent
CALLINAN LAWRIE , Private Bag 7, KEW VIC 3101
- (56) Prior Art Documents
WO 93/19670
- (57) Claim

1. A method of deconvolution of response data obtained from transmission of a MLS (including similar and variant sequences as hereinbefore defined), comprising multiplying each sample of incoming data in turn by the elements of a recovery sequence as the sample is received and immediately adding the product into corresponding positions in a reconstruction buffer, so that the samples of incoming data are deconvoluted and stored in the reconstruction buffer as they are received.

2. A method according to claim 1, applied to recording of biological signals which constitute said incoming data and which are obtained as response data due to stimulation by a MLS.

3. A method according to claim 2 and for testing hearing, wherein the response data are evoked otoacoustic emissions.

OPI DATE 21/11/94 APPLN. ID 65736/94
AOJP DATE 05/01/95 PCT NUMBER PCT/GB94/00862



AU9465736

CT)

(51) International Patent Classification 5 : G06F 15/336		A1	(11) International Publication Number: WO 94/25925 (43) International Publication Date: 10 November 1994 (10.11.94)
(21) International Application Number: PCT/GB94/00862 (22) International Filing Date: 22 April 1994 (22.04.94) (30) Priority Data: 9308715.3 27 April 1993 (27.04.93) GB (71) Applicant (for all designated States except US): MEDICAL RESEARCH COUNCIL [GB/GB]; 20 Park Crescent, London W1N 4AL (GB). (72) Inventors; and (75) Inventors/Applicants (for US only): THORNTON, Arthur, Roger, David [GB/GB]; 105 Deacon Road, Bitterne, Southampton, Hants SO2 8PT (GB). CHAMBERS, John, David [GB/GB]; 59 Osbourne Grove, Sherwood, Nottinghamshire NG5 2HE (GB). FOLKARD, Timothy, John [GB/GB]; 8 Enfield Street, Beeston, Nottinghamshire NG9 1AL (GB). (74) Agent: KEITH W. NASH & CO.; Pearl Assurance House, 90-92 Regent Street, Cambridge CB2 1DP (GB).		(81) Designated States: AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, ES, FI, GB, GE, HU, JP, KG, KP, KR, KZ, LK, LU, LV, MD, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TJ, TT, UA, US, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i> 674576	
(54) Title: DECONVOLUTION OF MLS RESPONSE DATA			
(57) Abstract A method of and apparatus for the deconvolution of response data obtained from transmission of a maximum length sequence (MLS), wherein the samples of incoming data are deconvoluted and stored in a reconstruction buffer as said samples are received. It is possible in this way to employ a reconstruction buffer shorter than that necessary to store the complete deconvoluted waveform.			

- 1 -

Title: Deconvolution of MLS Response Data**Field of the Invention**

This invention relates to a method of and apparatus for the deconvolution of response data obtained from transmission of maximum length sequences (MLS), similar sequences or variants of similar sequences. References to MLS hereinafter are intended to include such similar or variant sequences.

Background to the Invention

The applicants' prior International Patent Application No PCT/GB93/00639. relates to the use of a maximum length sequence technique to record otoacoustic emissions. One field of application of the present invention is the deconvolution of response data in the technique proposed in the prior patent application. However, the present invention is not restricted to that field of application.

For signals that have to be extracted from noise, a conventionally applied method, as described in the prior patent application, is to stimulate with an MLS, record the response or incoming signal generated by that MLS and, using sampling techniques, store the received waveform in a memory buffer whose length is equal to the length of the MLS multiplied by the number of ADC samples that are required per MLS entry in order to avoid aliasing errors.

- 2 -

Another MLS is immediately delivered and its received waveform added to that already in the memory buffer. This process is continued until sufficient received waveforms have been added to improve the signal-to-noise ratio to an adequate degree.

The original response, or input signal, then has to be recovered by a deconvolution technique. One example of a suitable technique, proposed in the prior patent application, is as follows.

Let L equal the length of the MLS. The MLS represents stimuli that will be presented at a specified rate. If the minimum time between presentations of stimuli, corresponding to the elements in the MLS, is such that n samples are required to characterise the response or incoming signal then that period is known as a slice of the MLS and n is the number of samples in one slice. Once the responses to a set of MLSs have been recorded, as described above, then the resultant waveform is contained in a raw data buffer whose length is equal to nL . A reconstruction memory buffer, of equal length to the raw data buffer, is zeroed. The recovery sequence is then generated and comprises a 1 for every instance for which the MLS contains a 1 and a -1 for every occurrence of a zero in the original MLS. The data in the raw data buffer are then each multiplied by the first element in the recovery sequence. The product is added to the reconstruction buffer. The data in the raw data buffer are then rotated left by one slice. They are then each multiplied by the second element in the recovery sequence and again added to the reconstruction buffer. This operation is continued until all the elements in the recovery sequence have been used. The reconstruction

- 3 -

buffer will then contain the original waveform deconvolved from the MLS. The procedure is illustrated in Figure 1 of the accompanying drawings.

The disadvantages of the above-described deconvolution technique may be illustrated by an example using evoked otoacoustic emissions recorded with an MLS.

1. As the response is being averaged from noise then it is useful to be able to avoid the noisiest periods (excess noise) during the recording of the signal. Typically this is done by rejecting, that is not adding to the average, any periods of the response where the waveform recorded from the subject or patient exceeds a certain limit. However, as the stimulus is also recorded then, using conventional techniques, the stimulus period has to be excluded from this rejection criterion. With the MLS technique and the overlapping of stimuli and responses, it is not possible sensibly to apply the described rejection methods. This is a problem with using the above-described deconvolution technique for the MLS. The signal-to-noise ratio is poorer than it might otherwise be because periods of high noise activity have to be included in the average.

2. Without the use of a very large amount of processing power, no deconvolved waveform can be obtained until the entire averaging process is complete because errors will occur if the averaging procedure is interrupted in order to effect the deconvolution. This is because MLS samples will be missed, leading to errors in the reconstruction of the original waveform.

The Invention

According to one aspect of the present invention, there is provided a method of deconvolution of response data obtained from transmission of a MLS (including similar and variant sequences as hereinbefore defined), comprising multiplying
5 each sample of incoming data in turn by the elements of a recovery sequence as the sample is received and immediately adding the product into corresponding positions in a reconstruction buffer, so that the samples of incoming data are deconvoluted and stored in the reconstruction buffer as they are received.

10 For completeness, it should be mentioned that an MLS is a quasi-random binary sequence with strictly defined mathematical properties. Possible variants are Legendre sequences, M-pulse sequences and De Bruijn sequences.

15 In a development of the method and apparatus of the invention, means are provided to reject immediately any one or more reconstructed MLSs which are contaminated by noise in excess of a predetermined limit such that they should not be added to the average being collected in the reconstruction buffer.



- 5 -

The invention has the advantage that a raw data buffer memory, equal in size to the length of the MLS, is no longer required. As a result, it is possible to achieve substantial saving in the time required to carry out deconvolution and reconstruction of the original signal.

Furthermore, the reconstruction buffer need only be as large as the time window required to observe the final waveform. This means that said buffer can, in general, be very much shorter than the MLS. Thus, it is possible to transmit very long MLSs without the requirement for large buffers.

One field of application of the invention is in the recording of biological signals obtained as response data due to stimulation by a MLS. For example, in the testing of hearing, a MLS may be used to record Evoked Otoacoustic Emissions (EOAs) which constitutes the incoming response data. The original signal can then be recovered from this response data by the deconvolution method and apparatus in accordance with the invention.

The apparatus in accordance with the invention may comprise a digital signal processing board, a computer and controlling software.

Thus, according to another aspect of the invention, there is provided a digital signal processing board and a computer programmed to carry out deconvolution of the MLS by the method hereinbefore defined.

- 6 -

Description of Embodiment

The method and apparatus of the invention are exemplified in the following description, making reference to accompanying Figures 1 to ~~4~~⁵, in which:-

Figure 1 illustrates a prior deconvolution technique;

Figure 2 illustrates the deconvolution technique in accordance with the invention;

Figure 3 is a flow diagram illustrating the technique in accordance with the invention, when carried out using a recovery window shorter than the MLS; and

Figure 4 is a table of values applicable to the flow diagram of Figure 3 in one typical case.

A prior deconvolution technique is illustrated in Figure 1. For an MLS of length L , the slices B_0 to B_{L-1} of the incoming signal are averaged and stored in a raw data buffer. The values in the data buffer are then multiplied by the first element in the recovery sequence (M_0 - M_{N-1}). The data from the raw data buffer are then added to the previously zeroed reconstruction buffer. The data in the raw data buffer are then rotated one place to the left, multiplied by the second element in the recovery sequence and added to the contents of the reconstruction buffer. This process is repeated until all distinct rotations of the raw data buffer have been completed (a total of L additions and $L-1$ rotations).

From Figure 1 it can be seen that samples initially stored in slice B_n are ultimately added to A_m after multiplying

- 7 -

by M_{n-m} . In accordance with the invention, it has been realised that, instead of adding raw samples to B and then shifting, multiplying and adding into A, it is possible to simply multiply each raw sample in turn by the appropriate element M of the recovery sequence and add it directly to the corresponding positions in A. This will build up the reconstructed average directly from the incoming samples. This process is illustrated in Figure 2.

This approach in accordance with the invention has several advantages:-

1. With each incoming sample being dealt with on arrival there is no need to store the raw data samples for subsequent processing; the large raw data buffer B (of size nL) is thus dispensed with.
2. Only the R slices of A that span the required reconstruction window need to be stored, thus the memory required for the reconstruction buffer is reduced from nL to nR.
3. The number of operations performed on each sample is dependent on R rather than L, thus the processing load is reduced.
4. With both memory and processing requirements reduced, one can more readily use double-buffering techniques to generate a separate reconstruction for each presentation of the MLS sequence. Each reconstructed response can then be compared with a template and rejected if any value is too large. Acceptable responses can then be added into a summation buffer. The direct reconstruction approach thus makes real-time rejection of excess noise periods

- 8 -

possible. This is a marked practical advantage for taking measurements where the time to record is limited primarily by the noise contamination on the signal.

Another way of explaining the technique in accordance with the invention is as follows.

Figure 3 is a flow diagram illustrating an application of the technique in accordance with the invention when the recovery window or reconstruction buffer is shorter than the MLSS. The logical operations employed in the technique will be readily understood from the flow diagram, when this is read in conjunction with the following definitions:-

L	Length of MLS
n	Number of samples in one slice of the MLS
m	Number of slices in the recovery window
p	Number of samples in MLS ($p=n*L$)
x	ADC samples of the input ($i=1$ to p)
q	Number of samples in the recovery window ($q=m*n$)
W(j)	Recovery window ($j=1$ to q)
R(k)	Recovery sequence [$= +1$ or -1] ($k=1$ to L)
a	Number of slices to start of recovery window
z	Number of MLS needed in average
mptr	Main recovery sequence pointer
rprr	Secondary recovery sequence pointer
wptr	Recording window pointer

Further understanding will be gained from Figure 4, which is a table of flow diagram values for one particular case in which one MLS has a length 3 and the recovery window

- 9 -

starts after one slice and has a duration of two slices.

In principle, the illustrated technique, which may be referred to as MLS recovery "on the fly", is as follows:-

Consider an MLS of length 3. This will have slices M1, M2 and M3 as shown in [A].

For the MLS in which M1=M2=1 and M3=0, the rotation and multiplication needed for conventional recovery are shown in [B].

[A]

M1	M2	M3
----	----	----

[B]
$$\begin{array}{rrrrrrrr} 1 & 1 & 0 & 1 & 1 & 1 & 0 \\ 1 & 0 & 1 & \times & 1 & = & 1 & 0 & 1 \\ 0 & 1 & 1 & -1 & 0 & -1 & -1 \\ \hline \text{SUM:} & 2 & 0 & 0 \end{array}$$

In general, the multiplication and rotations are shown in [C]. It may not always be the case that the required recovery window will start at the start of the MLS and a case in which the recovery window has a delay of one slice from the start of the MLS is shown in [D].

[C]

M1	M2	M3
M2	M3	M1
-M3	-M1	-M2

[D]

M1	M2	M3
M2	M3	M1
-M3	-M1	-M2

Now consider the same example as shown in [D] but with the individual ADC samples taken into the picture. The MLS is 1 1 0, with a recovery sequence $R(k) = 1 \ 1 \ -1$. Let there be three ADC samples per slice ($n=3$). The ADC samples in one MLS will be denoted by the letters a to i.

The MLS and the samples are shown in [E] and the matrix of samples that will make up the recovered waveform for the delayed recovery window, are shown in [F].

[E]

1	1	0						
a	b	c	d	e	f	g	h	i

[F]

a	b	c	d	e	f	g	h	i
d	e	f	g	h	i	a	b	c
-g	-h	-i	-a	-b	-c	-d	-e	-f

The flow diagram and the accompanying table shows how the on-the-fly recovery algorithm can be used to obtain the desired, recovered waveform.

- 10 -

The invention is preferably realised using a digital signal processing board and a conventional computer. The software may be realised in any suitable language or code and, as an example, details of the routines FILLER and CHECKER which have been written in "C", to give an example of the coding involved, are as follows:-

```
/*=====
FILLER - collect & deal with one sample
```

In actual use this would be an interrupt service routine, dealing with each incoming sample. For each input sample a stimulus sample may be output, depending on the value of the current MLS entry.

```
=====*/
```

```
void filler () {
    float samp;                // current input sample
    int n;                     // -> MLS entries
    int * mpr;                 // -> MLS entry for current slice

    mpr = &mls[sliceno];      // -> MLS entry for current slice

    if (*mpr==1)               // entry indicates stimulus?
        outsamp (outbuff[sampno]); // yes- output stimulus sample

    samp = getsamp();          // get input sample

    for (n=0; n<rlen; n+=slen) { // for each recon. slice
        currbuff [n+sampno] += samp * *mpr; // add/subtract to current buffer
        mpr--;                 // move back through MLS...
        if (mpr < mls)         // with wrap-around to end of MLS
            mpr = &mls[mlslen-1];
    }

    sampno++;                  // count samples done within slice
    if (sampno < slen)         // return if still within slice
        return;

    // start of new slice:-
    sampno = 0;                // reset sample number
    sliceno++;                 // bump slice counter
    if (sliceno < mlslen)      // return if not at end of MLS
        return;

    // end of MLS:-
    sliceno = 0;               // reset slice counter
    toggle = !toggle;         // switch reconstruction buffers
    currbuff = toggle ? recon1 : recon2; // -> next buffer
    mlspt = &mls[sliceno];    // Reset mls pointer to start of MLS
}
```

- 11 -

```

/*=====
CHECKER - process completed reconstructions

Checks to see if a buffer has completed being filled.  If so, checks buffer
values against a rejection template and, if no values are too large, adds each
value into a summation buffer.
=====*/

int checker () {
    int i;
    int ok;

    if (currbuff == prevbuff)           // has 'filler' finished a recon.?
        return (0);                     // no- wait some more

                                        // yes- 'prevbuff' points at it...
    ok = 1;
    for (i=0; i<rlen; i++)               // check each value against template
        if (fabs(prevbuff[i]) > reject[i]) {
            ok = 0;
            break;
        }

    if (ok) {                            // all values passed-
        for (i=0; i<rlen; i++)           // add to summation buffer
            summation[i] += prevbuff[i]; // count good reconstructions
        mlsdone++;
    }

    for (i=0; i<rlen; i++)               // re-zero buff for next reconstruction
        prevbuff[i] = 0;

    prevbuff = currbuff;                 // update pointer
}

/*=====
MAIN

In practice 'checker' would be written to loop until enough good
reconstructions had been gathered, and would be continuously interrupted
by 'filler'.  We simulate this here by placing both calls in a loop.
=====*/

main (int argc, char * argv[]) {
    int i, n, n_needed;

    n_needed = 10;                       // ask for 10 reconstructions
    mls_init ();                          // initialise variables, buffers, etc

    do {
        filler ();
        checker ();
    } while (mlsdone < n_needed);

    for (i=0; i<rlen; i+=4) {             // dump results
        for (n=0; n<4; n++)
            printf ("%15.4f", summation[i+n]);
        printf ("\n");
    }
}

```

The claims defining the invention are as follows:-

1. A method of deconvolution of response data obtained from transmission of a MLS (including similar and variant sequences as hereinbefore defined), comprising multiplying each sample of incoming data in turn by the elements of a recovery sequence as the sample is received and immediately adding the product into corresponding positions in a reconstruction buffer, so that the samples of incoming data are deconvoluted and stored in the reconstruction buffer as they are received.

2. A method according to claim 1, applied to recording of biological signals which constitute said incoming data and which are obtained as response data due to stimulation by a MLS.

3. A method according to claim 2 and for testing hearing, wherein the response data are evoked otoacoustic emissions.

4. Apparatus for deconvolution of response, data obtained from transmission of a MLS (including similar and variant sequences as hereinbefore defined), comprising a reconstruction buffer and means for deconvoluting each sample of incoming data as it arrives and storing the deconvoluted sample in the reconstruction buffer, wherein the means for deconvoluting include means for multiplying each sample in turn by the elements of the recovery sequence as the sample is received and immediately adding the product into corresponding portions in the reconstruction buffer.

5. Apparatus according to claim 4, in which means are provided to reject immediately any one or more reconstructed MLSs which are contaminated by



noise in excess of a predetermined limit such that they should not be added to the average being collected in the reconstruction buffer.

6. Apparatus according to claim 4 or 5, in which the reconstruction buffer is only as large as the time window required to observe a final waveform of the stored data.

7. Apparatus according to any of claims 4 to 6, which comprises a digital signal processing board, a computer and controlling software.

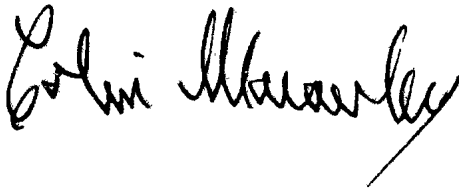
8. Apparatus comprising a digital signal processing board and a computer programmed to carry out deconvolution of the MLS by the method of any of claims 1 to 3.

DATED this 30th day of October, 1996.

MEDICAL RESEARCH COUNCIL

By their Patent Attorneys:

CALLINAN LAWRIE



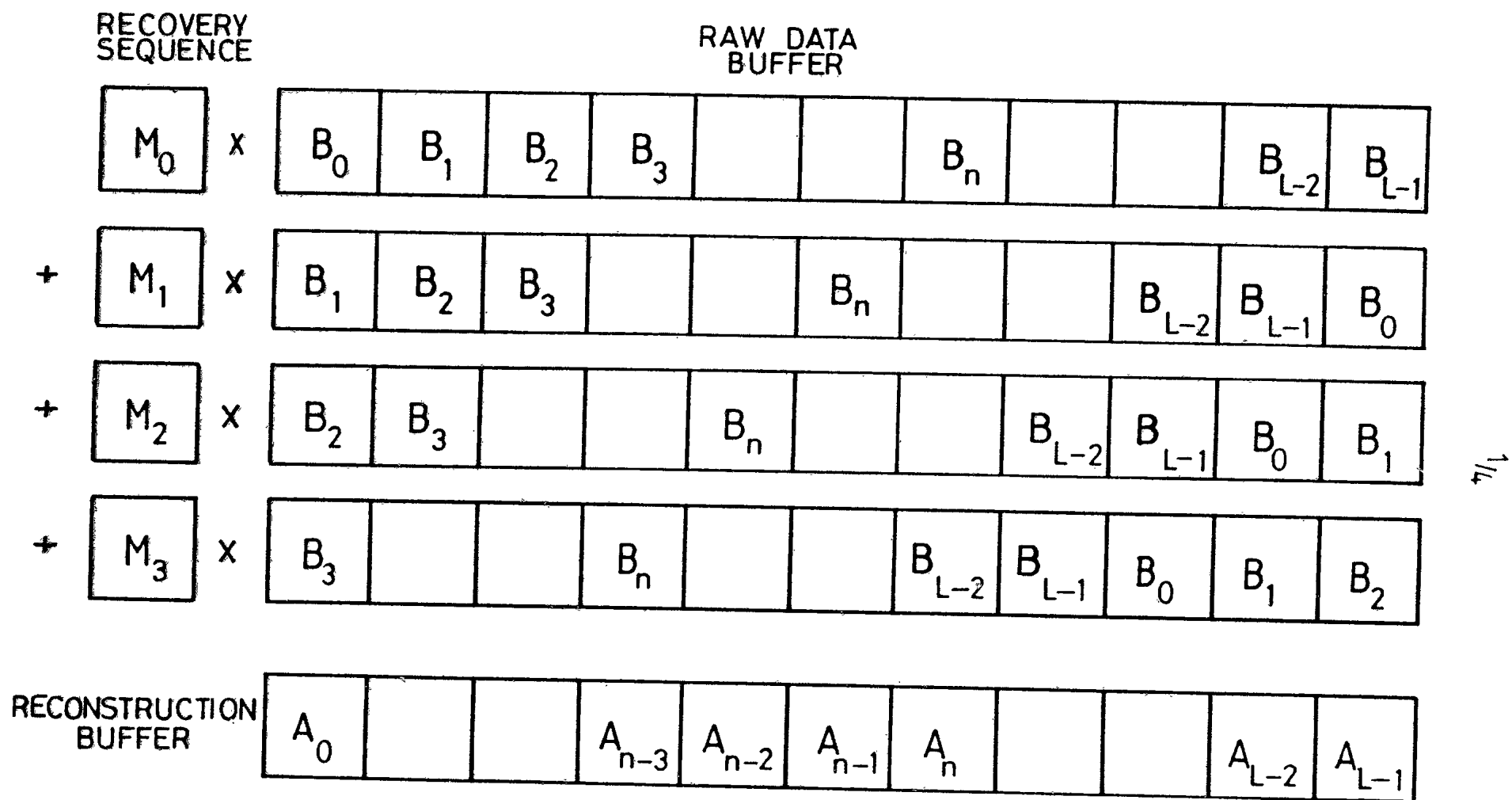


Fig. 1 PRIOR TECHNIQUE

65736/14

2/4

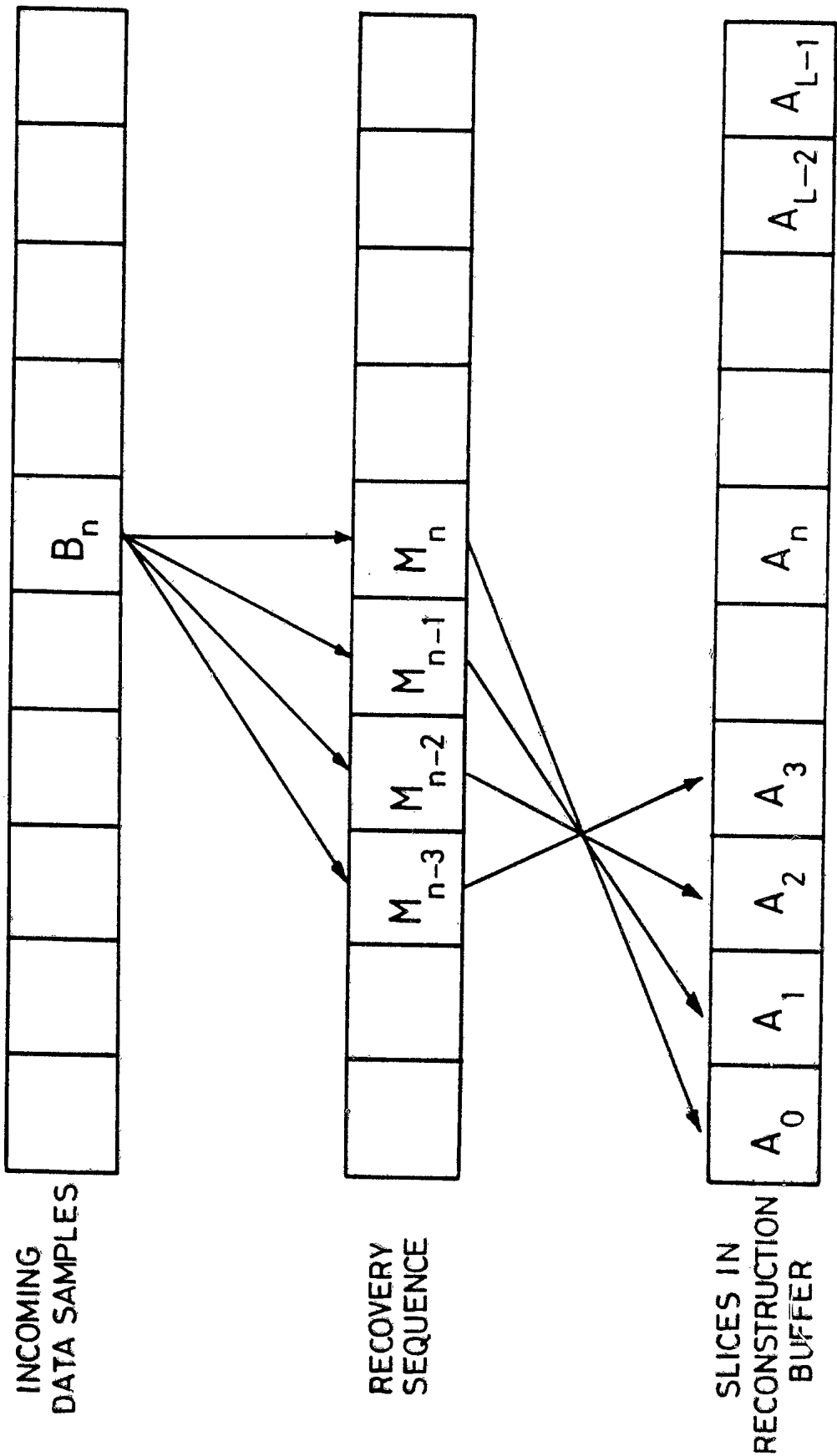
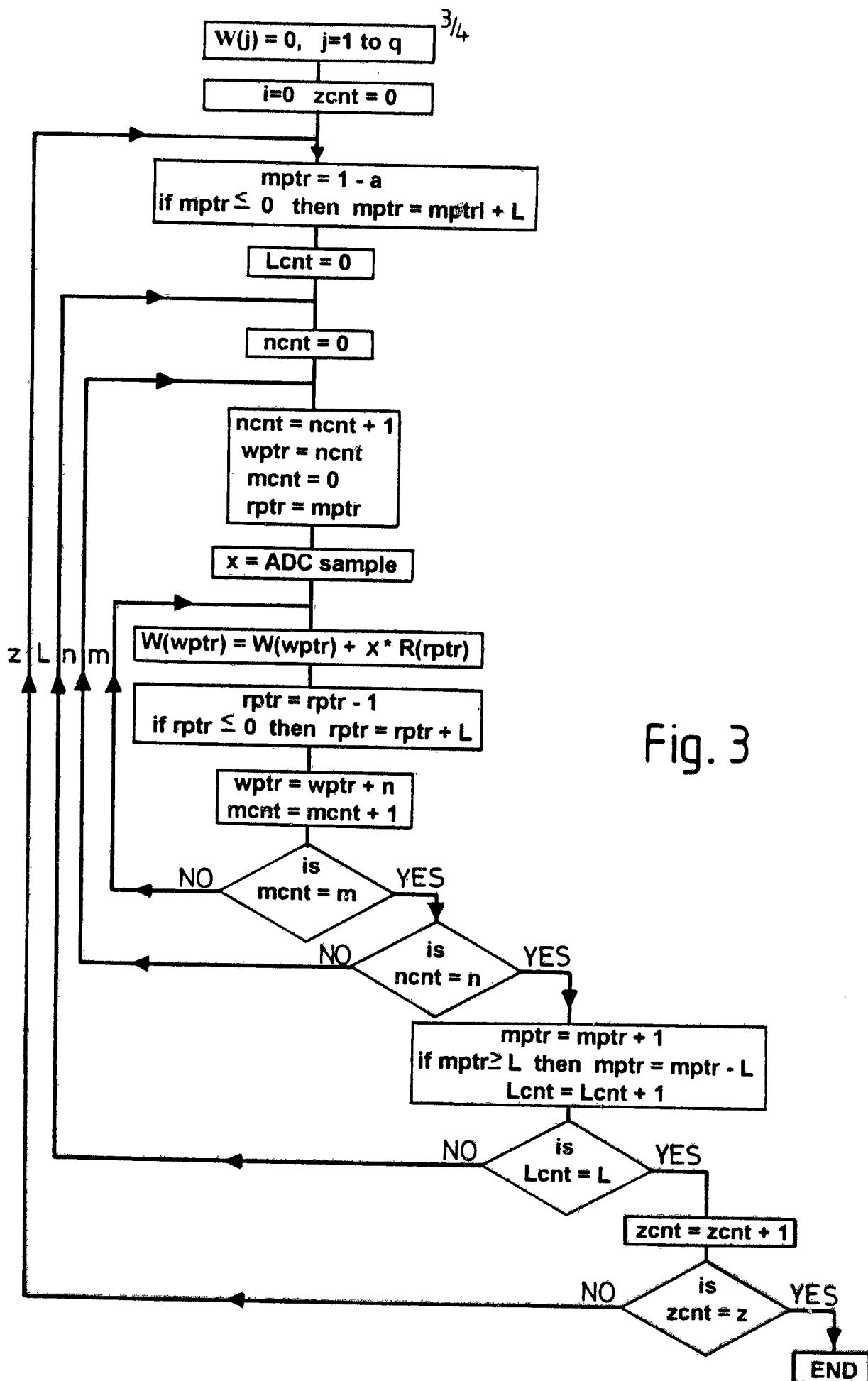


Fig. 2



4/4
TABLE

L=3
n=3
m=2
a=1
z=1

rp_{tr}: 1 2 3
R(rp_{tr}): 1 1 -1

INIT. CONDS.	sample	Lcnt	ncnt	mcnt	mptr	rp _{tr}	wptr	RECOVERY WINDOW					
	0	0	0	0	3	3	1	0	0	0	0	0	0
	1	0	1	1	3	3	1	-a					
		0	1	2	3	2	4				a		
	2	0	2	1	3	3	2	-b					
		0	2	2	3	2	5					b	
	3	0	3	1	3	3	3		-c				
		0	3	2	3	2	6						c
	4	1	1	1	1	1	1	d					
		1	1	2	1	3	4				-d		
	5	1	2	1	1	1	2		e				
		1	2	2	1	3	5					-e	
	6	1	3	1	1	1	3			f			
		1	3	2	1	3	6						-f
	7	2	1	1	2	2	1	g					
		2	1	2	2	1	4				g		
	8	2	2	1	2	2	2		h				
		2	2	2	2	1	5					h	
	9	2	3	1	2	2	3			i			
		2	3	2	2	1	6						i

Fig. 4

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/GB 94/00862

A. CLASSIFICATION OF SUBJECT MATTER

IPC 5 G06F15/336

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 5 A61B G06F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MEDICAL AND BIOLOGICAL ENGINEERING AND COMPUTING, vol.30, January 1992, STEVENAGE GB pages 32 - 40 CHAN ET AL. 'Measurement of human BAERs by the maximum length sequence technique' see page 32, left column, line 1 - page 36, right column, line 12; figures 1-5 --- -/--	1,3,8



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *A* document member of the same patent family

Date of the actual completion of the international search

23 August 1994

Date of mailing of the international search report

27 09 94

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Chen, A

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JOURNAL OF THE ACOUSTICAL SOCIETY OF AMERICA, vol.87, no.4, April 1990, NEW YORK US pages 1656 - 1664 BURKARD ET AL. 'A comparison of maximum length and Legendre sequences for the derivation of brain-stem auditory-evoked responses at rapid rates of stimulation' see page 1656, left column, line 1 - page 1658, right column, line 11; figures 1,2 ---	1-3,8
A	PROCEEDINGS OF THE ANNUAL INTERNATIONAL CONFERENCE OF THE IEEE ENGINEERING IN MEDICINE AND BIOLOGY SOCIETY, vol.11, November 1989, US pages 1289 - 1290 YING SHI ET AL. 'The use of M pulse sequences in the study of nonlinearities in the brainstem auditory evoked response' see the whole document -----	1