EDV in Medizin und Biologie 14 (3–4), 98–108, ISSN 0300-8282 © Verlag Eugen Ulmer GmbH & Co., Stuttgart; Gustav Fischer Verlag KG, Stuttgart

## Statistical analysis of single-case experimental designs:

Conditional equivalence of the general-linear-model approach of GLASS, WILLSON & GOTTMAN with the intervention model of BOX & TIAO

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#### Summary

In single case diagnostics the researcher has often to evaluate the influence of his intervention. If the data are at the level of the interval-scales the impact assessment can be pursued by the GLASS, WILLSON & GOTTMAN (1975) approach, which can be regarded as a mixture of the general linear model and time series modelling of the error component, or by the intervention model of BOX & TIAO (1975), which is a descendant of BOX & JENKINS' transfer model (1970).

In clinical research only the former method is used for impact assessment. The latter model is up to now not widely known. Even in statistical literature there is no unified treatment<sup>1</sup> of both approaches. We want to show that they are equivalent under certain conditions. But in spite of this, researchers are advised to abandon the GLASS, WILLSON & GOTTMAN method in favor of the intervention model because of its greater elegance and practicability. This means that the danger of misspecifying the intervention effects is far more negligible in the intervention model than in the GLASS, WILLSON & GOTTMAN method.

#### Zusammenfassung

In der Einzelfalldiagnostik sollen oft Interventionseffekte evaluiert werden. Sind die Daten intervallskaliert, kann das nach dem Ansatz von GLASS, WILLSON & GOTTMANN (1975) erfolgen. Er beinhaltet die Modellierung des Effektes nach dem allgemeinen linearen Modell und die des Fehlers nach dem ARIMA-Modell von BOX & JENKINS. Eine andere Möglichkeit der Interventionsevaluation bietet das Interventionsmodell von BOX & TIAO (1975), das sich aus dem Transfermodell von BOX & JENKINS (1970) herleitet.

In der klinischen Forschung wird meist nur die erste Methode verwendet. Das zweite Modell ist bis jetzt nicht sehr bekannt. Sogar in der statistischen Literatur gibt es keine einheitliche Darstellung<sup>2</sup> beider Ansätze. Wir wollen zeigen, daß sie unter

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bestimmten Bedingungen äquivalent sind. Dennoch sollte in Untersuchungen das Interventionsmodell wegen seiner größeren Eleganz und Praktikabilität vorgezogen werden. Außerdem ist im Interventionsmodell die Gefahr von Fehlspezifikationen des Interventionseffektes kleiner als im Ansatz von GLASS, WILLSON & GOTTMANN.

# 1. The general linear model and many-case experimental designs

The analysis of cross-sectional data arising from N>1-experiments with the general linear model is well known (WOT-TAWA, 1974; MOOSBRUGGER, 1978; TIMM, 1975; BOCK, 1975). The structure of the model is in the univariate case

(1.1) 
$$_{N}y_{1} = {}_{N}\underline{X}_{M}\beta_{1} + {}_{N}\underline{\varepsilon}_{1}$$

where: N = number of subjects

M = number of variables

$$_{N}y_{1} = N \times 1$$
 vector of the dependent variable

 ${}_{N}\overline{X}_{M} = N \times M$  matrix of M predictors

- (= design matrix)
- $_{M}\beta_{1} = M \times 1$  vector of parameters
- $N\overline{\epsilon}_1 = N \times 1$  vector of error variables

To obtain the best linear unbiased (blue) estimators of  $\beta$  it is usual to formulate a »weak« set of assumptions (1.2a–1.2c):

(1.2a) 
$$Y_i = \underline{x}_i^{\prime}\beta + \varepsilon_i$$

where the  $\epsilon_i \ (i{=}1{,}{\ldots}{,}N)$  are independent random variables with

(1.2b) 
$$E(\varepsilon_i) = 0$$
 or  $E(\varepsilon) = 0$ 

and variance

1.2c) 
$$\operatorname{var}(\varepsilon_i) = \sigma_{\varepsilon}^2$$
 or  $\operatorname{E}(\underline{\varepsilon} \ \underline{\varepsilon}') = \sigma_{\varepsilon}^2 \cdot \underline{\mathrm{I}}$ 

Two additional assumptions are often not stated explicitly, though they will be important in this context:

(1.3a) <u>X</u> must not contain measurement errors (GOLD-BERGER, 1973). Especially <u>X</u> must not contain estimators of parameters.

and

# (1.3b) $\underline{X}$ has to be known apriori and must not contain unknown parameters.

As an example we want to write down the familiar t-test with the two hypotheses  $H_0$ :  $\mu_A = \mu_B$  and  $H_1$ :  $\mu_A \neq \mu_B$  in terms of (1.1). The full model representing the alternative  $H_1$  hypothesis can be written

<sup>&</sup>lt;sup>1</sup> JENKINS (1979) does not mention the GLM approach. Only KEESER (1979, p. 268) gives a short informal hint concerning possible parallelism in both approaches. Even the new book of GOTTMAN (1981, p. 365ff) treats both approaches separately without any reference to their partial identity.

<sup>&</sup>lt;sup>2</sup> JENKINS (1979) erwähnt nicht den Ansatz von GLASS, WILLSON & GOTTMAN. Nur KEESER (1979, S. 268) gibt einen kurzen informellen Hinweis auf mögliche Parallelen zwischen den beiden Modellen. Sogar das neue Buch von GOTTMAN (1981, S. 365 ff.) behandelt beide Ansätze separat, ohne einen Hinweis auf ihre partielle Identität zu geben.

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 $y = \underline{X}_R \beta_R + \underline{\varepsilon}_R$ 

 $\hat{\beta}_{R} = (\underline{X}_{R}, \underline{X}_{R})^{-1} \underline{X}_{R}, \underline{y}$ 

$$\begin{array}{c} 1 \\ 2 \\ 3 \\ group A \\ \vdots \\ (1.4a) \\ 1 \\ group B \\ \vdots \\ N_{B} \end{array} \begin{bmatrix} Y_{11} \\ Y_{12} \\ Y_{13} \\ \vdots \\ Y_{1N_{A}} \\ Y_{21} \\ Y_{22} \\ Y_{23} \\ \vdots \\ Y_{2N_{B}} \end{bmatrix} = \begin{bmatrix} 1 & 0 \\ 1 & 0 \\ 1 & 0 \\ \vdots & \vdots \\ 1 & 0 \\ \vdots & \vdots \\ 1 & 0 \\ \vdots & \vdots \\ 1 & 1 \\ 1 & 1 \\ \vdots & \vdots \\ 1 & 1 \end{bmatrix} + \left( \begin{bmatrix} L_{F} \\ \beta_{F} \end{bmatrix} \right) \\ + \left( \begin{bmatrix} 1 \\ 0 \\ 1 \\ 0 \\ \vdots \\ 1 \\ 1 \\ 1 \\ 1 \\ \vdots \\ 1 \\ 1 \end{bmatrix} \right) \\ or$$

 $X_F = designmatrix$ 

$$\beta_{\rm F}$$
 = vector of parameters in full model

The ordinary least squares (OLS) estimator is

(1.5) 
$$\hat{\beta}_{\rm F} = (\underline{X}_{\rm F}^{2} \underline{X}_{\rm F})^{-1} \underline{X}_{\rm F}^{2} y$$

and the estimate of the residual vector

(1.6) $\underline{\hat{e}}_{F} = \underline{e}_{F} = y - \underline{X}_{F}\hat{\beta}_{F}$ 

Under H<sub>0</sub> we expect the equality of population means:  $\mu_A = \mu_B$ . The equivalent hypothesis for our model (1.4) is:  $\beta_F = 0$ . To check the hypothesis, we insert  $\beta_F = 0$  in (1.4) and formulate the reduced model

(1.7b) The OLS-estimator is (1.8)

or

(1.9)

811F ε<sub>12F</sub>

E<sub>13F</sub>

EINAF

 $\epsilon_{21F}$ 

822F

E23F ;

E2NBF

:

$$\hat{\underline{e}}_{R} = \underline{e}_{R} = \underline{y} - \underline{X}_{R}\underline{\beta}_{I}$$

Now we can test H<sub>0</sub> with the F-ratio

(1.10) 
$$F_{df_{1},df_{2}} = \frac{(\underline{e}_{R}^{2}\underline{e}_{R} - \underline{e}_{F}^{2}\underline{e}_{F})/(p_{F} - p_{R})}{(\underline{e}_{F}^{2}\underline{e}_{F})/(N - p_{F})}$$
where: N = number of subjects
$$p_{F} = number of linear independent$$
columns in  $\underline{X}_{F}$  (here: 2)
$$p_{R} = number of linear independent$$
columns in  $\underline{X}_{R}$  (here: 1)
$$df_{1} = p_{F} - p_{R}$$

$$df_{2} = N - p_{F}$$

If we take the square root of (1.10) we get the t-ratio with  $df = N - p_F$ .

### 2. The general linear model and single-case experimental designs

In the single-case experimental design, which is not a crosssectional but a longitudinal design, one subject is observed under various experimental conditions (CHASSAN, 1972). The impact of the experimental conditions should be assessed by statistical methods (BARLOW & HERSEN, 1973; HERSON & BARLOW, 1976; KAZDIN, 1976; FICHTER, 1979; TYLER & BROWN, 1968).

One of the most familiar designs is the A1B1A2B2-design. The observation time is partitioned into four intervals. Two intervals (A1,A2) are without experimental intervention. A1 is sometimes called »base line«. B1 represents the first intervention phase. An example for an even more complicated  $A_1B_1C_1B_2C_2$ -experiment is demonstrated in Figure 1.

Without loss of generality we want to restrict our attention to the simple AB-design. The impact of the intervention can be analyzed according the general linear model along (1.1)-(1.9) if the statistical assumptions (1.2a)-(1.3b) are met



Fig. 1. Data from an experiment examining the effect of feedback on the cating behaviour of a patient with anorexia nervosa (Patient 4). (Fig. 3, p. 283, from: Agras, W. S., Barlow, D. H., CHAPIN, H. N., ABEL, G. G., and LEI-TENBERG, H. Behaviour modification of anorexia nervosa. Archives of General Psychiatry, 1974, 30, 279-286. Reproduced by permission.)

by the data. In that case the full model is (1.4) and the reduced model is (1.7).  $N_A$  and  $N_B$  are now the number of time points, where measurements are taken in phase A and B so that  $N_A + N_B = T$ . Whereas in a cross-sectional design it is assumed that the  $\varepsilon_i$  and  $Y_i$  are independent, in a time-series experiment it is possible to test this assumption. We have to prove that the time-series of estimated residuals  $\hat{\varepsilon}_t = e_t$  is sampled from a time-discrete white-noise process  $a_t$ . For »diagnostic checks applied to residuals« we refer to Box & JENKINS (1976, p. 287–299).

If the residuals  $\boldsymbol{\epsilon}_i$  are dependent and not a white-noise process, then

(2.1) 
$$E(\underline{\varepsilon} \ \underline{\varepsilon}') = \sigma^2 \cdot \underline{\Omega} \neq \sigma^2 \cdot \underline{I}$$

and the estimators are no longer blue. But more important for impact assessment is the fact, that the F-value (1.10) is »too large« or »too small«, depending on the correlation structure of the  $\varepsilon$ -process (HIBBS, 1974). Nonsignificant effects could appear to be significant and vice versa!

In the case of (2.1) it is at least in principle possible to estimate the parametervector  $\underline{\beta}$  with the generalized least squares (GLS)-method (AITKEN, 1935). The linear model (1.1), (1.4) and (1.7) must be transformed, so that the new residuals  $\varepsilon_1^*$  will be independent and will follow a white-noise process. We have to look for a T × T transformation matrix A, with

(2.2a) 
$$\underline{\mathbf{A}} \cdot \underline{\mathbf{\Omega}} \cdot \underline{\mathbf{A}}' = \underline{\mathbf{I}}$$

(2.2b) 
$$\frac{\underline{A}}{\underline{y}^*} = \underline{AX\beta} + \underline{A} \underline{\epsilon}$$
$$\frac{\underline{y}^*}{\underline{y}^*} = \underline{X}^*\underline{\beta} + \underline{\epsilon}^*$$
with:  $y^* = \underline{A} y$ ,  $\underline{X}^* = \underline{A} \cdot \underline{X}$ ,  $\underline{\epsilon}^* = \underline{A} \underline{\epsilon}$ 

and

(2.3) 
$$E(\underline{\varepsilon}^{*}\underline{\varepsilon}^{*'}) = E(\underline{A} \ \underline{\varepsilon} \ \underline{\varepsilon}^{'}\underline{A}^{'}) = \underline{A} \cdot E(\underline{\varepsilon} \ \underline{\varepsilon}^{'}) \cdot \underline{A}^{'}$$
$$= \underline{A} \cdot \sigma_{\varepsilon}^{2} \underline{\Omega} \cdot \underline{A}^{'} = \sigma_{\varepsilon}^{2} \cdot \underline{A} \cdot \underline{\Omega} \cdot \underline{A}^{'} = \sigma_{\varepsilon}^{2} \cdot \underline{I}$$

Because of the positive-definiteness of  $\underline{\Omega}$ , <u>A</u> is a triangular matrix (HIBBS, 1974; REVENSTORFF & KEESER, 1979). The GLS-estimator of  $\beta$  is

(2.4) 
$$\hat{\underline{\beta}} = (\underline{X}^{*}\underline{X}^{*})^{-1}\underline{X}^{*}\underline{y}^{*} \\ = (\underline{X}^{*}\underline{A}^{*}\underline{A}^{*}\underline{X})^{-1}\underline{X}^{*}\underline{A}^{*}\underline{A}^{*}\underline{y} = (\underline{X}^{*}\underline{\Omega}^{-1}\underline{X})^{-1}\underline{X}^{*}\underline{\Omega}^{-1}\underline{y}$$

The GLS-model is equivalent to transformation (2.2) and successive OLS. But usually the autocovariance matrix  $\underline{\Omega}$  of the residuals is unknown and has to be estimated. The estimation is impossible, if all parameters in  $\underline{\Omega}$  are free and if the time-series is finite. So we have to assume some simplifying structure in  $\underline{\Omega}$ , so that the number of unknown parameters decreases sharply. It is the merit of GLASS, WILLSON & GOTTMAN (1975) who provided us with a wide variety of new design matrices  $\underline{X}^*$  for various  $\varepsilon_t$ -processes.

#### 3. The approach of GLASS, WILLSON & GOTTMAN

On the basis of an early article of BOX & TIAO (1965), GLASS, WILLSON & GOTTMAN (1975) expanded the GLS-approach covering a wide variety of intervention designs and autocorrelation structures of  $Y_t$  and  $\varepsilon_t$ .

3.1 If e.g. the raw data follow a moving-average-process of order 1 (= ARIMA (0,0,1) in the BOX & JENKINS terminology), we expect the data for the preintervention phase A to be construed in accordance with the model

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(3.1a) 
$$\begin{aligned} Y_t &= L - \theta_1 a_{t-1} + a_t = f(L, \theta_1, a_{t-1}, a_t) \\ \text{with} \quad \underline{a} \sim N(\underline{0}, \sigma^2 \cdot \underline{I}) \end{aligned}$$

and for the intervention-phase B along the model

(3.1b) 
$$Y_{t} = (L + \beta) - \theta_{1}a_{t-1} + a_{t} = f(L,\beta,\theta_{1},a_{t-1},a_{t})$$
  
with  $-1 < \theta_{1} < +1$ 

which can be combined with the linear model (full model):

	1 2	1 2 3	$\begin{bmatrix} Y_{11} \\ Y_{12} \end{bmatrix}$		1	0 0	$\begin{bmatrix} L_F \\ \beta_F \end{bmatrix}$		$\begin{bmatrix} 0 \\ a_1 \end{bmatrix}$	$\begin{bmatrix} a_1 \\ a_2 \end{bmatrix}$	$\begin{bmatrix} -\theta_1 \\ 1 \end{bmatrix}$
phase A	3	3	Y <sub>12</sub> Y <sub>13</sub>		1	ŏ			a24	a <sub>3</sub>	Γ., Έ
(e.g.: baseline)	-21	:	:	1	ł	:			:	:	
		T <sub>A</sub>	Y <sub>1TA</sub>		1	0		38			
(3.2a)		<del></del>	10000	1	1.00			+			
	•2	1	Y <sub>21</sub>		1	1			- 63	а I	
	•	1 2 3	Y <sub>22</sub>	- 5	1	1					
phase B	•	3	Y <sub>23</sub>		1	1			10		
(e.g.: inter-	:	÷	:			÷			1	:	
vention)	Т	$T_B$	$Y_{2T_B}$		1	1			a <sub>T-1</sub>	a <sub>T</sub>	
			-	<b>1</b> 10			5			~	, —
										<u></u>	
or											

(3.2b)

and:

with: 
$$\varepsilon_t = -\theta_1 a_{t-1} + a_t = (1 - \theta_1 B) a_{t-1}$$

 $= X \beta + \varepsilon$ 

where B is the backshift operator

$$a_t = a_{t-1}$$
 and in general  $B^k a_t = a_{t-k}$ 

$$Da_t = a_{t-1}$$
 and in general  $Da_t =$ 

(3.3)  $E(\underline{\varepsilon} \ \underline{\varepsilon}') = \sigma_{\varepsilon}^2 \cdot \underline{\Omega} \neq \sigma_{\varepsilon}^2 \cdot \underline{I}$ 

E

The dependence of the residuals (3.3) can be seen very quickly in (3.2a). 
$$\epsilon_{t-1}$$
 and  $\epsilon_t$  share a common term, which is  $a_{t-1}$ .

Now we want to determine the matrices  $\sigma_{\epsilon}^2 \cdot \underline{\Omega}$ , <u>A</u> and <u>X</u><sup>\*</sup>. The variance of the correlated residuals is

(3.4) 
$$\operatorname{var}(\varepsilon_t) = \operatorname{var}(-\theta_1 a_{t-1}) + \operatorname{var}(a_t) = (1 + \theta_1^2)\sigma_a^2$$

and the covariance of lag k=1

(3.5) 
$$\operatorname{cov}(\varepsilon_{t},\varepsilon_{t-1}) = \operatorname{cov}(-\theta_{1}a_{t-1} + a_{t}, -\theta_{1}a_{t-2} + a_{t-1}) \\ = -\theta_{1}\sigma_{a}^{2}$$

so the autocorrelation of both time series  $Y_k$  and  $\epsilon_k$  with their lagged counterparts  $Y_{t-k}$  and  $\epsilon_{t-k}$  is

(3.6) 
$$\varrho_{k} = \begin{cases} \frac{-\theta_{1}}{1+\theta_{1}^{2}} & k = 1\\ 0 & k > 1 \end{cases}$$

The autocovariance matrix of the residuals can be written as the symmetric matrix

$$(3.7) \ \mathrm{E}(\underline{\varepsilon}\underline{\varepsilon}') = \begin{bmatrix} 1 & \ddots & \ddots & \ddots & \vdots \\ -\theta_{1} & 1 & \ddots & \ddots & \vdots \\ 3 \\ \vdots \\ T \end{bmatrix} \begin{bmatrix} 1 & \ddots & \ddots & \ddots & \vdots \\ -\theta_{1} & 1 \cdot \ddots & \ddots & \vdots \\ 0 & \ddots & 1 \cdot \theta_{1}^{2} & 1 \cdot \ddots & \ddots \\ \vdots & \ddots & \ddots & \ddots & \vdots \\ 0 & \ddots & \ddots & 0 & \frac{\cdot -\theta_{1}}{1 + \theta_{1}^{2}} & 1 \end{bmatrix} \sigma_{a} (1 + \theta_{1}^{2})$$

(

(3.7) shows that only in the case of  $\theta_1 = 0$ , the residuals are independent and the familiar t-test is justified to test the intervention hypothesis H<sub>0</sub>:  $\beta_F = 0$ .

The transformation matrix  $\underline{A}$  is (REVENSTORFF & KEESER, 1979):

(3.8) 
$$\underline{A} = \begin{bmatrix} 1 & & & \\ 2 & \\ 3 & \\ \vdots & \\ T & \\ \end{bmatrix} \begin{bmatrix} 1 & & & \\ \theta_1 & 1 & & \\ \theta_1^2 & \theta_1 & 1 & \\ \vdots & \vdots & \vdots & \ddots & \\ \theta_1^{T-1} & \theta_1^{T-2} & \theta_1^{T-3} & & 1 \\ 1 & 2 & 3 & \cdots & T \end{bmatrix}$$

For the transformed designmatrix  $\underline{X}^* = \underline{A} \underline{X}$  we get:

and as the transformed vector of the dependent variable (= timeseries  $Y_t$ ):

(3.10a)

$$\begin{bmatrix} \mathbf{Y}_{11}^{*} \\ \mathbf{Y}_{12}^{*} \\ \mathbf{Y}_{13}^{*} \\ \vdots \\ \mathbf{Y}_{T}^{*} \end{bmatrix} = \begin{bmatrix} \mathbf{Y}_{11} \\ \mathbf{Y}_{12} + \theta_1 \mathbf{Y}_{11} \\ \mathbf{Y}_{13} + \theta_1 \mathbf{Y}_{12} + \theta_1^2 \mathbf{Y}_{11} \\ \vdots \\ \mathbf{Y}_{T} + \theta_1 \mathbf{Y}_{T-1} + \dots + \theta_1^{T-1} \mathbf{Y}_{11} \end{bmatrix}$$
$$= \begin{bmatrix} \mathbf{Y}_{11}^{*} \\ \mathbf{Y}_{12} + \theta_1 \mathbf{Y}_{11} \\ \mathbf{Y}_{13} + \theta_1 \mathbf{Y}_{12}^{*} \\ \vdots \\ \mathbf{Y}_{T} + \theta_1 \mathbf{Y}_{T-1}^{*} \end{bmatrix}$$
(2.10b)

(3.10b) <u>y</u><sup>\*</sup> = <u>A y</u>

The transformed residual factor  $\underline{\epsilon}^* = \underline{A} \ \underline{\epsilon}$  takes the simple form

(3.11)  $\underline{\varepsilon}^* = \underline{A} \underline{\varepsilon} = \underline{a}$ As a result we get the transformed linear model (2.2) with independent errors

(3.12) 
$$\underline{y}^* = \underline{X}^*\underline{\beta} + \underline{\varepsilon}^* = \underline{X}^*\underline{\beta} + \underline{a}$$
  
with  $\underline{a} \sim N(\underline{0}, q_a^2; \underline{I})$ 

If we estimate the full and reduced models, we can test the intervention hypothesis with the familiar F-ratio (1.10). Higher order moving average processes (ARIMA(0,0,q)-models) can be transformed in a similar fashion.

3.2 Now the general transformation method will be demonstrated for the general nonstationary ARIMA(p,d,q)-model. In literature three alternative formulations of the same ARIMA-process (version I, II, III) are used.

3.2.1 Three formulations for the general ARIMA(p,d,q)-model: We postulate that the preintervention data are construed in accordance with the ARIMA(p,d,q)-model (HIBBS, 1977, p. 140; MÖBUS & NAGL 1983)

3.13a)  
Version I  
3.13b) 
$$\begin{cases} \Phi_{p}(B)(1-B)^{d}Y_{t} = \theta_{o} + \theta_{q}(B)a_{t} \\ Y_{t} = \frac{\theta_{o} + \theta_{q}(B)a_{t}}{\Phi_{p}(B)(1-B)^{d}} = \frac{\theta_{q}(B)}{\Phi_{p}(B)(1-B)^{d}}a_{t} + \frac{\theta_{o}}{\Phi_{p}(B)(1-B)^{d}} \\ \text{where:} \quad a) \quad \Phi_{o} (B) \text{ is the autoregressive operator:} \end{cases}$$

b) 
$$\theta_q$$
 (B) is the autoregressive operator:  
 $\Phi_p$  (B) =  $(1 - \Phi_1 B - \ldots - \Phi_p B^p)$   
b)  $\theta_q$  (B) is the moving average operator:  
 $\theta_q$  (B) =  $(1 - \theta_1 B - \ldots - \theta_q B^q)$   
c)  $(1-B)^d$  is the difference operator:  
e.g.:  $(1-B)^2 Y_t = (1 - 2B + B^2) Y_t$   
 $= Y_t - 2Y_{t-1} + Y_{t-2}$   
which is exactly taking differences of differences  
e.g.:  $(1-B)^2 Y_t = \nabla^2 Y_t = (\nabla Y_t - \nabla Y_{t-1})$ 

- $= (Y_t Y_{t-1}) (Y_{t-1} Y_{t-2})$ d)  $\theta_o$  is an unknown constant
- e) and the inverse of the differenceoperator is e.g. for (1-B) defined as:

$$\frac{1}{(1-B)} = (1-B)^{-1} = (1+B+B^2+...) = \sum_{k=0}^{\infty} B^k$$
  
as can be shown by multiplication:  
 $(1 + B + B^2 + ...) (1-B) = 1$   
f) and the inverse of e.g.  $\Phi_1$  (B) is  $(1 - \Phi_1 B)^{-1}$   
 $= (1 + \Phi_1 B + \Phi_1^2 B^2 + ...) = \sum_{k=0}^{\infty} \Phi_1^k B^k$   
as can be shown too:  
 $(1 + \Phi_1 B + \Phi_1^2 B^2 + ...) (1 - \Phi_1 B) = 1$ 

Coefficients of a general  $\Phi_p^{-1}(B)$  an  $(1-B)^{-d}$  can be obtained by equating coefficients.

Various authors use a second model formulation. If we put the differencing operator  $(1-B)^d$  on the left side of the equality sign of (3.13b), we can express the nonstationary model in differences  $w_t = (1-B)^d Y_t$  (JENKINS, 1977, p. 98):

(3.14a)  
Version II  
(3.14b) 
$$\begin{cases} (1-B)^{d}Y_{t} = \frac{\theta_{q}(B)}{\Phi_{p}(B)} a_{t} + \frac{\theta_{o}}{\Phi_{p}(B)} \\ w_{t} = \frac{\Phi_{q}(B)}{\Phi_{p}(B)} a_{t} + \mu_{w} \end{cases}$$

where: 
$$\mu_w = E(w_t) = \frac{\theta_o}{\Phi_p(B)} = \frac{\text{mean of the d-th}}{\text{differences}}$$

A third representation is preferred by GLASS, WILLSON & GOTTMAN (1975). They choose

$$L = \frac{\mu_{w}}{(1-B)^{d}} = \frac{\theta_{o}}{\Phi_{p}(B)(1-B)^{d}}$$

and put L on the left side of (3.13b), so that we may write

$$\begin{array}{ll} (3.15a) \\ \text{Version III} \\ (3.15b) \end{array} \quad \begin{cases} \mathbf{Y}_t - \mathbf{L} = \frac{\theta_q(\mathbf{B})}{\Phi_p(\mathbf{B}) (1-\mathbf{B})^d} \mathbf{a}_t \\ \Phi_p(\mathbf{B})(1-\mathbf{B})^d (\mathbf{Y}_t - \mathbf{L}) = \theta_q(\mathbf{B}) \mathbf{a}_t \end{cases}$$

(3.15) is an infinite polynominal in B for d>0 and/or p>0. This means, that the deviation  $(Y_t - L)$  can be expressed as an infinite moving average process. This is called the »random shock« form (BOX & JENKINS, 1976, p. 95 ff).

(3.16a) 
$$Y_t - L = \frac{\theta_q(B)}{\Phi_p(B)(1-B)^d} a_t = \psi(B)a_t$$

where: 
$$\psi(\mathbf{B}) = \psi_0 + \psi_1 \mathbf{B} + \psi_2 \mathbf{B}^2 + \dots$$
  
with  $\psi_0 = 1$ 

or

W

W

(3.16b) 
$$Y_t = L + \sum_{k=1}^{n} \psi_k a_{t-k} + a_t = L$$

To get the unknown  $\psi$ -weights we multiply (3.16a) on both sides with  $(1-B)^d \Phi_p(B)$ .

(3.17) 
$$\Phi_{p}(B)(1-B)^{d}(Y_{t}-L) = \Phi_{p}(B)(1-B)^{d}\psi(B)a_{t}$$

Following (3.15b), the left side of (3.17) is  $\theta_0(B)a_t$ . So we get

(3.18) 
$$\theta_{q}(B)a_{t} = \Phi_{p}(B)(1-B)^{d}\psi(B)a_{t} = \phi(B)\psi(B)a_{t}$$

The equation of operators is

(3.19a)  $\theta_{0}(B) = \varphi(B)\psi(B)$ 

> $\phi(B)$  is the general autoregressive operator where: (Box & JENKINS, 1976, p. 95):

$$\varphi(\mathbf{B}) = \varphi_{p}(\mathbf{B})(1-\mathbf{B})^{d} = \varphi_{o} - \varphi_{1}\mathbf{B} - \varphi_{2}\mathbf{B}^{2} - \dots - \varphi_{p+d}\mathbf{B}^{p+d}$$

with  $\varphi_o = 1$ 

and can be written explicitly as:

(3.19b) 
$$(1-\theta_1B-...-\theta_qB^q) = (1-\varphi_1B-...-\varphi_{p+d}B^{p+d})(1+\psi_1B+\psi_2B^2+...)$$

Therefore, the  $\psi$ -weights can be obtained by equating coefficients in B on the left side of (3.19) to the coefficients in B of the right side.

Theoretically there are countably infinite coefficients  $\psi_k$ . GLASS et al. (1975, p. 152) argue, that the values of unobserved time series  $Y_k$ ,  $a_k$  ( $k \leq 0$ ) should be set to their expected value, so that only  $\psi_1, \ldots, \psi_{t-1}$  have to be derived  $(\psi_o$ is set to 1).

(3.20) 
$$Y_t = L' + \sum_{k=0}^{t-1} \psi_k a_{t-k} + a_t$$

The derivation of the  $\psi$ -weights is shown explicitly in appendix A, because the treatment of GLASS et al. (1975, p. 153-154) is confusing.

3.2.2. Transformation of the  $\psi$ -weight form of the intervention-model to the general linear model

The linear intervention-model (full model) in  $\psi$ -weight form is

$$\begin{array}{c} \text{phase } \mathbf{A} \begin{bmatrix} \mathbf{Y}_{11} \\ \mathbf{Y}_{12} \\ \mathbf{Y}_{13} \\ \vdots \\ \mathbf{Y}_{1T_{A}} \\ \hline \mathbf{Y}_{21} \\ \mathbf{Y}_{22} \\ \mathbf{Y}_{23} \\ \vdots \\ \mathbf{Y}_{2T_{B}} \end{bmatrix} = \begin{bmatrix} 1 & 0 \\ 1 & 0 \\ 1 & 0 \\ \vdots & \vdots \\ 1 & 0 \\ \vdots & \vdots \\ 1 & 0 \\ \hline \vdots & \vdots \\ 1 & 0 \\ \hline \vdots & \vdots \\ 1 & 1 \\ \vdots & \vdots \\ 1 & 1 \\ \vdots & \vdots \\ \mathbf{Y}_{2T_{B}} \end{bmatrix} = \begin{bmatrix} 1 & 0 \\ 1 & 0 \\ \vdots & \vdots \\ 1 & 0 \\ \hline \vdots & \vdots \\ \vdots \\ \vdots \\ 1 & 1 \\ \vdots \\ 1 & 1 \\ \vdots \\ 1 & 1 \\ \vdots \\ \mathbf{Y}_{T-1} \psi_{T-2} \psi_{T-3} \dots \psi_{1} \end{bmatrix} \begin{bmatrix} \mathbf{a}_{1} \\ \mathbf{a}_{2} \\ \mathbf{a}_{3} \\ \vdots \\ \vdots \\ \vdots \\ \vdots \\ \mathbf{a}_{T} \end{bmatrix} \\ (3.21b) \quad \underline{\mathbf{Y}} = \underline{\mathbf{X}} \underbrace{\boldsymbol{\beta}} \\ \mathbf{y} = \underline{\mathbf{Y}} \underbrace{\boldsymbol{a}} \end{bmatrix} + \underbrace{\mathbf{W}} \underline{\mathbf{a}}$$

Some authors prefer instead of using matrix  $\underline{\Psi}$ , the formulation with the operator  $\psi(B)$ 

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(3.21c) 
$$\underline{Y} = \underline{X} \beta + \psi(\underline{B})\underline{a}$$

 $+\sum_{k=0}^{\infty}\psi_k a_{t-k}$ 

(3.21d) 
$$Y_t = f_t(L', \beta_1, ..., \beta_{s+1}) + \psi(B)a$$

The residuals ε<sub>1</sub> are not independent

(3.22) 
$$E(\underline{\varepsilon} \ \underline{\varepsilon}') \neq \sigma_{\varepsilon}^2 \cdot \underline{I}$$

which is due to the overlap of common terms in  $\underline{\varepsilon}$ .

(3.23) 
$$\underline{\varepsilon} = \begin{cases} \psi_0 a_1 \\ \psi_1 a_1 + \psi_0 a_2 \\ \psi_2 a_1 + \psi_1 a_2 + \psi_0 a_3 \\ \psi_3 a_1 + \psi_2 a_2 + \psi_1 a_3 + \psi_0 a_4 \end{cases}$$

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The variance-covariance matrix of the correlated residuals  $\varepsilon_t$ is:  $\operatorname{cov}(\underline{\varepsilon} \ \underline{\varepsilon}') = \sigma_a^2 \cdot \underline{\Omega} =$ 

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$$\sigma_{a}^{2} \cdot k \begin{bmatrix} \psi_{o}^{2} & \psi_{o}\psi_{1} & \psi_{o}\psi_{2} & \dots & \psi_{o}\psi_{l-1} & \dots & \psi_{o}\eta \\ \hline \psi_{o}\psi_{1} & \psi_{o}^{2} + \psi_{1}^{2} & \sum_{i=1}^{2}\psi_{2-i}\psi_{3-i} & \dots & \sum_{i=1}^{2}\psi_{2-i}\psi_{l-i} & \dots & \eta_{i=1}^{2}\psi_{2-i} \\ \hline \psi_{o}\psi_{2} & \sum_{i=1}^{2}\psi_{2-i}\psi_{3-i} & \sum_{i=1}^{3}\psi_{i-1} & \dots & \sum_{i=1}^{3}\psi_{3-i}\psi_{l-i} & \dots & \eta_{i=1}^{3}\psi_{3-i} \\ \hline \psi_{o}\psi_{3} & \sum_{i=1}^{2}\psi_{2-i}\psi_{4-i} & \sum_{i=1}^{3}\psi_{3-i}\psi_{4-i} & \dots & \sum_{i=1}^{4}\psi_{4-i}\psi_{1-i} & \dots & \eta_{i=1}^{3}\psi_{4-i} \\ \hline \vdots & \vdots & \vdots & \vdots & \ddots & \vdots \\ \hline \psi_{o}\psi_{k-1} & \sum_{i=1}^{2}\psi_{2-i}\psi_{k-i} & \sum_{i=1}^{3}\psi_{3-i}\psi_{k-i} & \dots & \sum_{i=1}^{\min\{k,1\}}\psi_{k-i}\psi_{k-i} & \dots & \eta_{i=1}^{k}\psi_{k-i} \\ \hline \vdots & \vdots & \vdots & \vdots & \vdots & \vdots \\ \hline \psi_{o}\eta & \eta_{i=1}^{2}\psi_{2-i} & \eta_{i=1}^{3}\psi_{3-i} & \dots & \eta_{i=1}^{1}\psi_{1-i} & \dots & \sum_{i=1}^{\infty}\psi_{i-1} \\ \hline where: & \eta = \lim_{j\to\infty}\psi_{j} \end{bmatrix}$$

The intervention-model with correlated residuals (3.21) can be transformed to one with uncorrelated residuals. This can be done either by multiplying (3.21) by  $\Psi^{-1}$ 

(3.25) 
$$\underline{\Psi}^{-1} = \sum_{i=0}^{\infty} (\underline{I} - \underline{\Psi})^{i}$$
  
(3.26a) 
$$\underline{\Psi}^{-1}\underline{Y} = \underline{\Psi}^{-1}\underline{X} \underline{\beta} + \underline{a}$$
  
(3.26b) 
$$\underline{Y}^{*} = \underline{X}^{*}\underline{\beta} + \underline{a}$$

or by using the recursive relationships

(3.27a) 
$$Y_{t}^{*} = Y_{t} - \sum_{j=1}^{t-1} \psi_{j} Y_{t-j}^{*}$$
  
(3.27b)  $x_{tk}^{*} = x_{tk} - \sum_{j=1}^{t-1} \psi_{j} x_{t-j,k}^{*}$   
(3.27c)  $a_{t} = \varepsilon_{t} - \sum_{j=1}^{t-1} \psi_{j} a_{t-j}$ 

For T=5 time points we get the following matrix  $\Psi^{-1}$ , where  $T_A=2$  and  $T_B=3$ :  $(3.28) \quad {}_{5}\Psi_{5}^{-1} =$ 

The burden of computations is taken by computer programs, distributed by BOWER, PADIA & GLASS (1974) and GOTTMAN (1981) and will thus not be demonstrated here.

#### 4. The intervention-model-approach of Box & TIAO

BOX & TIAO (1975) developed their intervention model on the basis of BOX & JENKINS' transfer function model (BOX & JENKINS, 1970). Their intervention model is given by:

(4.1)  $\delta(B)Y_t = \Omega(B)I_t + \delta(B)\varepsilon_t$ 

where  $\varepsilon_t$  is assumed to be an ordinary ARIMA(p,d,q)-model according (3.13b)

(4.2a) 
$$\epsilon_{t} = \frac{\theta_{o}}{(1-B)^{d} \Phi_{p}(B)} + \frac{\theta_{q}(B)}{(1-B)^{d} \Phi_{p}(B)} a_{t}$$
with  $L = \frac{\theta_{o}}{(1-B)^{d} \Phi_{p}(B)}$ 

According (3.16a) we obtain

(4.2b) 
$$\varepsilon_t = L + \psi(B)a_t$$

Substituting equation (4.2b) into (4.1), we get the form:

(4.3a) 
$$\delta(B)Y_t = \Omega(B)I_t + \delta(B) \cdot \{L + \psi(B)a_t\}$$

or equivalently

(4.3b) 
$$Y_{t} = \underbrace{\frac{\Omega(B)}{\delta(B)} \cdot I_{t}}_{\mathscr{Y}_{t}} + L + \psi(B)a_{t}$$

or equivalently

(4.3c) 
$$Y_t - L = \begin{cases} effects \\ of intervention \\ Y_t \end{cases} + \begin{cases} noncontrollable \\ effects \\ N_t \end{cases}$$

 $\mathcal{Y}_t$  is that part of time-dependent data, which is controlled by the experimenter. The intervention variable I<sub>t</sub> is at apriori determined time points set to '1' (=»on«) or '0' (=»off«). The aim of intervention analysis is the splitting of the timeseries  $\mathcal{Y}_t$  and N<sub>t</sub> and the estimation of the parameters of the  $\mathcal{Y}_t$ -process. These parameters are effect-parameters which can be tested.

»The function  $\mathscr{Y}_t$  represents the additional effects of the intervention over the noise. In particular, when N<sub>t</sub> is nonstationary, large changes could occur in the output even with no intervention. Fitting the model can make it possible to distinguish between what can and what cannot be explained by the noise.« (Box & TIAO, 1975, p. 72), HIBBS (1975) calls the transfer function model

(4.4a) 
$$\mathscr{Y}_{t} = \frac{\Omega(B)}{\delta(B)} \cdot I_{t}$$

or equivalently

(4.4b) 
$$\delta(B)\mathcal{Y}_t = \Omega(B)I_t$$

or equivalently

(4.4c) 
$$\delta(B) \mathcal{Y}_t = \omega(B) B^b I_t$$

or equivalently

or equivalently

(4.4e) 
$$\mathcal{Y}_{t} - \delta_{1} \mathcal{Y}_{t-1} - \dots - \delta_{r} \mathcal{Y}_{t-r}$$
  
=  $\omega_{o} I_{t-b} - \omega_{1} I_{t-b-1} - \dots - \omega_{s} I_{t-b-s}$ 

the »general intervention effects model«. The indices reflect the »memory« of the intervention component. If we have a nonintervention phase A and an intervention phase B, the intervention variable  $I_t$  takes the values

If there is an abrupt change in level of  $Y_t$  without time-delay, we use the intervention model of order zero

(4.5) 
$$\mathcal{Y}_t = \omega_0 I_t$$
 step change without time-delay  
(Fig. 2a)

If there is a step change with time delay, we use

(4.6) 
$$\mathcal{Y}_{t} = \omega_{o} I_{t-b}$$
 step change with time delay b  
(Fig. 2b)

In the case of a nonstationary  $N_t$ , the level of  $Y_t$  shows a different reaction.

Is the effect  $\mathcal{Y}_t$  changing slowly, we use the transfer function of order one

$$\begin{array}{ll} (4.7a) \quad \mathscr{Y}_{t} = \delta_{1} \quad \mathscr{Y}_{t-1} + \omega_{o} I_{t-b} \\ \text{or} \\ (4.7b) \quad (1 - \delta_{1}B) \quad \mathscr{Y}_{t} = \omega_{o} I_{t-b} \\ \text{or} \\ (4.7c) \quad \mathscr{Y}_{t} = \frac{\omega_{o}}{(1 - \delta_{1}B)} \cdot I_{t-b} \end{array} \right\} \begin{array}{l} \text{ramp change} \\ \text{with} \\ \text{time delay b} \\ (\text{Fig. 2c}) \end{array}$$

Is  $\delta_1 = 0$  and b = 0, we get again the step change (Fig. 2d) and in the case of  $\delta_1 = 1$ , we get the model

(4.8) 
$$\mathcal{Y}_{t} = \frac{1}{(1-B)} \cdot I_{t-b}$$
 nondamped increase (Fig. 2e)

which is not stable.

The modelling of various other intervention effects is shown in MÖBUS & NAGL (1983).

The stability of the intervention model (4.4) is guaranteed, if the roots  $B_1,...,B_r$ , of the characteristic equation

 $\delta(\mathbf{B}) = (1 - \delta_1 \mathbf{B} - \delta_2 \mathbf{B}^2 - \dots - \delta_r \mathbf{B}^r) = 0$ 

satisfy the conditions  $|B_j| > 1$  for j = 1,...,r. For the intervention model of order two

$$(1 - \delta_1 \mathbf{B} - \delta_2 \mathbf{B}^2) \mathcal{U}_t = \omega_0 \mathbf{I}_t$$

the roots B<sub>i</sub> of the polynomial

$$(1 - \delta_1 \mathbf{B} - \delta_2 \mathbf{B}^2) = 0$$

have to be of absolute value  $|B_i| > 1$ 

with 
$$B_j = \frac{\delta_1 \pm \sqrt{\delta_1^2 + 4\delta_2}}{2\delta_2}$$

The conditions  $|\mathbf{B}_{j}| > 1$  can only be met, if the parameters lie in these regions:

$$\begin{array}{c} -1 < \delta_2 < +1 \\ \delta_1 + \delta_2 < +1 \\ \delta_2 - \delta_1 < +1 \end{array}$$

If the Intervention variable  $I_t$  is held constant  $I_t = 1$  for  $t > T_A$ ,  $\mathcal{Y}_t$  will reach the steady state:



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(4.9) 
$$E(\mathscr{Y}_{1}) = \mathscr{Y}_{\infty} = \frac{(\omega_{o} - \omega_{1} - \dots - \omega_{s})}{(1 - \delta_{1} - \dots - \delta_{r})}$$

BOX & TIAO (1975) and other authors distinguish between two forms of the indicator variable  $I_t$ :

the step input (4.10)  $I_t = S_t = \begin{cases} 0 \text{ for } t \leq T_A \\ 1 \text{ for } t > T_A \end{cases}$ 

and the pulse input

(4.11) 
$$I_t = P_t = \begin{cases} 0 \text{ for } t \neq T_A + 1 \\ 1 \text{ for } t = T_A + 1 \end{cases}$$

where the pulse input  $P_t$  can be derived by differencing the step input  $S_t$ 

(4.12)  $P_t = (1-B)S_t$ 

If there are k intervention variables acting simultaneously, the general intervention-effects-model can be extended similar to the statistical model of a multifactorial anova-model

(4.13) 
$$\mathcal{Y}_{t} = \sum_{j=1}^{k} \mathcal{Y}_{jt} = \sum_{j=1}^{k} \left\{ \frac{\Omega_{j}(B)}{\delta_{j}(B)} \right\} \cdot I_{jt}$$

For example it is reasonable to hypothesize two effects on the deterministic component of perceptual speed of a person in a drug experiment (Figure 3)

(4.14) 
$$\mathcal{Y}_{1} = \mathcal{Y}_{1t} + \mathcal{Y}_{2t} = f_{1}(I_{1t}) + f_{2}(I_{2t})$$
 see (5.1c)

where:  $I_{1t}$  is a step input: »no drug« vs. »drug«  $I_{2t}$  is a pulse input:  $(1-B)I_{1t}$ , which can be interpreted as the change of experimental conditions

The observable time series »perceptual speed« is contaminated by noncontrollable effects  $N_t$ , so that we get the time series model according to (4.3c)

$$Y_t - L = \mathcal{Y}_t + N_t$$

In practical data analysis it is advised to test a fixed sequence of intervention models. Hints can be found into CLEARY & HAY (1980) and MÖBUS & NAGL (1983).

### 5. The comparison of the GLS-approach of GLASS, WILL-SON & GOTTMAN with the Intervention-model of Box & TIAO

As shown in (3.21) the approach of GLASS et al. (1975) is given as a matrix equation by

(5.1a)  $\underline{Y} = \underline{X} \underline{\beta} + \underline{\Psi} \underline{a}$ 

with:  $\Psi$  = lower triangular matrix with  $\psi$ -weights

or in operator form either as

(5.1b)  $\underline{\mathbf{Y}} = \underline{\mathbf{X}} \underline{\boldsymbol{\beta}} + \boldsymbol{\psi}(\mathbf{B})\underline{\mathbf{a}}$ 

or for a single observation

(5.1c)  $Y_t = f_t(L', \beta_1, ..., \beta_{s+1}) + \psi(B)a_t$ 

with: L' = L only in the case of nontruncated white noise process  $a_t$ . If  $a_t$  is set to  $E(a_t)$  for t < 0, we have a truncated  $\psi$ -weight form and  $L' \neq L$ . In the further derivation it is not important wether we assume L = L'or  $L \neq L'$ .

Equation (5.1) represents the general form for testing s+1 effects of one or more interventions. The approach of Box & TIAO (1975) for testing s+1 effects of one intervention is (4.3)

$$(5.2) \qquad Y_t - L = \mathscr{Y}_t + \psi(B)a_t = \frac{\Omega(B)}{\delta(B)} \cdot I_t + \psi(B)a_t$$

If there are more than one interventions, with more than s+1 effects, we are allowed to extend the model as was done in (4.13). We want to show the equality of the functions  $f_t$  and  $L + \mathcal{Y}_t$  for fixed parameters  $\delta_i$  (i = 1,...,r) of  $\delta(B)$ :

(5.3a) 
$$L + \mathcal{Y}_t = f_t(L,\beta_1,...,\beta_{s+1})$$

or in matrix form

(5.3b) 
$$_{\mathrm{T}} \underline{L} + \underline{\mathscr{Y}}_{1} = _{\mathrm{T}} \underline{X}_{\mathrm{s}+2} \beta_{1}$$

Without loss of generality we suppose that we have to test s+1 effects of one intervention-variable I<sub>t</sub>. The general intervention-effect model  $\mathcal{Y}_t$  can be written in the form (4.4d) or

(5.4) 
$$\mathcal{Y}_{t} - \delta_1 \mathcal{Y}_{t-1} - \dots - \delta_r \mathcal{Y}_{t-r} = [I_{t-b} \ I_{t-b-1} \ \dots \ I_{t-b-s}] \underline{\omega}$$
  
where  $\underline{\omega}' = [\omega_o - \omega_1 \ \dots \ - \omega_s]$ 

or as

$$(5.5a) \mathcal{Y}_{t} = [I_{t-b} I_{t-b-1} \dots I_{t-b-s}] \underline{\omega} + \delta_{1} \mathcal{Y}_{t-1} + \dots + \delta_{r} \mathcal{Y}_{t-r}$$

In analogy to (5.4) we can write

$$(5.5b) \mathcal{Y}_{t-1} = [I_{t-b-1} \ I_{t-b-2} \dots I_{t-b-s-1}] \underline{\omega} + \delta_1 \mathcal{Y}_{t-2} + \dots + \delta_r \mathcal{Y}_{t-r-1}$$

$$(5.5c) \mathcal{Y}_{t-2} = [I_{t-b-2} I_{t-b-3} \dots I_{t-b-s-2}] \underline{\omega} + \delta_1 \mathcal{Y}_{t-3} + \dots + \delta_r \mathcal{Y}_{t-r-2}$$

$$(5.5d) \mathscr{Y}_{t-r} = [I_{t-b-r} I_{t-b-r-1} \dots I_{t-b-s-r}] \underline{\omega} + \delta_1 \mathscr{Y}_{t-r-1}$$

 $+ \dots + \delta_r \mathcal{Y}_{t-2r}$ 

In principle we need for the computation of  $\mathscr{Y}_t$  an infinite number of previous  $\mathscr{Y}s$ . But in practice  $\mathscr{Y}_t = 0$  for  $t \leq T_A + b$ , because  $I_t = 0$  for  $t \leq T_A$ .

As an example we'll compute  $\mathcal{Y}_t = h(I_t, I_{t-1}, ...)$  for  $t = T_A + b + 3$  and r = 2.

(5.6a) 
$$\mathscr{Y}_{T_{A}+b+3} = [I_{T_{A}+3} I_{T_{A}+2} I_{T_{A}+1} 0 \dots 0] \underline{\omega} + \delta_{1} \mathscr{Y}_{T_{A}+b+2} + \delta_{2} \mathscr{Y}_{T_{A}+b+1}$$
  
(5.6b)  $\mathscr{Y}_{T_{A}+b+2} = [I_{T_{A}+2} I_{T_{A}+1} 0 \quad 0 \dots 0] \underline{\omega} + \delta_{1} \mathscr{Y}_{T_{A}+b+1}$ 

$$(5.6c) \mathcal{Y}_{T_{A}+b+1} = [I_{T_{A}+1} 0 \quad 0 \quad 0 \dots 0] \underline{\omega}$$

$$(5.7a) \mathcal{Y}_{T_{A}+b+3} = [I_{T_{A}+3} I_{T_{A}+2} I_{T_{A}+1} 0 \dots 0] \underline{\omega}$$

$$+ \delta_{1} [I_{T_{A}+2} I_{T_{A}+1} 0 \quad 0 \dots 0] \underline{\omega}$$

$$+ \delta_{2} [I_{T_{A}+1} 0 \quad 0 \quad 0 \dots 0] \underline{\omega}$$

$$+ \delta_{1} [I_{T_{A}+2} I_{T_{A}+1} 0 \quad 0 \quad 0 \dots 0] \underline{\omega}$$

Collecting terms of  $\underline{\omega}$ :

(5.7b) 
$$\mathcal{Y}_{T_{A}+b+3} = \begin{bmatrix} I_{T_{A}+3} & I_{T_{A}+2} \\ + \delta_{1}I_{T_{A}+2} & \delta_{1}I_{T_{A}+2} \\ + (\delta_{1}^{2}+\delta_{2})I_{T_{A}+1} & \delta_{1}I_{T_{A}+1} \end{bmatrix} I_{T_{A}+1} \begin{bmatrix} 0 & \dots & 0 \\ \end{bmatrix} \underline{\omega}$$

Equation (5.7b) can be written shortly

(5.8)  $\mathcal{Y}_{T_A+b+3} = [g_{T_A+b+3,1} g_{T_A+b+3,2} g_{T_A+b+3,3} \dots g_{T_A+b+3,s+1}] \underline{\omega}$ In general we can express  $\mathcal{Y}_t$  similar to the example (5.8)

(5.9) 
$$\mathscr{Y}_{t} = [g_{t,1} g_{t,2} \dots g_{t,s+1}] \underline{\omega} \text{ for all } t > T_{A} + b$$
$$= g_{t}^{2} \underline{\omega}$$

The k-th component  $g_{t,k}$  (k = 1,...,s+1) of the vector  $\underline{g}_t$  can be computed as

(5.10) 
$$g_{t,k} = I_{t-b-k+1} + \sum_{l=1}^{N} \delta_l g_{t-l,k}$$
  
where  $g_{t,k} = 0$  for  $t \leq T_A + b + k - 1$ 

Now we define  $\beta_0 = L$ ,  $\beta_1 = \omega_0$  and  $\beta_j = -\omega_{j-1}$  (j = 2,...,s+1) and are able to write down the matrix equation (5.11)

(5.11a)  $_{\mathrm{T}}\underline{\mathbf{L}} + \underline{\mathscr{Y}}_{1} = {}_{\mathrm{T}}\underline{\mathbf{G}}_{s+1}\underline{\boldsymbol{\beta}}_{1}$ 

or explicitly

$$(5.11b) \begin{bmatrix} L + \mathcal{Y}_{1} \\ \vdots \\ L + \mathcal{Y}_{T_{A}+b} \\ \vdots \\ L + \mathcal{Y}_{T_{A}+b+2} \\ \vdots \\ L + \mathcal{Y}_{T} \end{bmatrix} = \begin{bmatrix} 1 & 0 & 0 & \dots & 0 \\ \vdots & \vdots & \vdots & \vdots \\ 1 & 0 & 0 & \dots & 0 \\ 1 & g_{T_{A}+b+1,1}0 & \dots & 0 \\ 1 & g_{T_{A}+b+2,1}g_{T_{A}+b+2,2} \dots & 0 \\ \vdots & \vdots & \vdots & \vdots \\ 1 & g_{T,1} & g_{T,2} & \dots & g_{T,s+1} \end{bmatrix} \begin{bmatrix} L \\ \beta_{1} \\ \vdots \\ \beta_{s+1} \end{bmatrix}$$

The elements  $g_{t,k}$  of the matrix <u>G</u> are fixed, because we assumed that the  $\delta$ 's are known a priori. In this case the design-matrix <u>X</u> can be set to <u>G</u> and (5.3) holds. Therefore the approaches of GLASS, WILLSON & GOTTMAN (1975) and BOX & TIAO (1975) are conditionally equivalent. This means that equivalence can be met only in the case of fixed  $\delta$  parameters.

When testing intervention hypothesis with the approach of BOX & TIAO (1975) we don't need any a priori information about the  $\delta$ 's. We estimate both the  $\delta$ 's and the  $\omega$ 's. If we try to evaluate the intervention-effect along GLASS et al., we have to know the estimates  $\hat{\delta}$ , to specify the design-matrix X. This is shown in the following example.

The transfer function model

(5.12) 
$$\mathscr{Y}_{t} = \frac{\omega_{o}}{1 - \delta_{1}B} \cdot P_{t}$$
  
with  $0 < \delta_{1} < 1$  and  $P_{t} = \begin{cases} 0 \text{ for } t \neq T_{A} + 1\\ 1 \text{ for } t = T_{A} + 1 \end{cases}$ 

describes an abrupt but temporary intervention effect. It follows from equation (5.12)

$$(5.13) \quad \mathcal{Y}_{t} = \delta_1 \mathcal{Y}_{t-1} + \omega_0 P_t$$

and we get

(5.14) 
$$\mathcal{Y}_{t} = \begin{cases} 0 & \text{for } t \leq T_{A} \\ \delta_{1}^{t-T_{A}-1} \cdot \omega_{o} & \text{for } t > T_{A} \end{cases}$$

The pattern of effect may be called a »decaying spike«. The parameter  $\delta_1$  is to be interpreted as the momentary rate of decay. When  $\delta_1$  is large, say  $\delta_1 = 0.9$ , the effect persists for a long period of time (1.0, 0.9, 0.81, ...). If  $\delta_1 = 0.5$  the decay is faster (1.0, 0.5, 0.25, ...). Testing the same hypothesis with the approach of GLASS et al. (1975) we have to know a priori the exact value of  $\delta_1$  in the design-matrix X:



This example has shown that the statistical hypothesis in the approach of GLASS, WILLSON & GOTTMAN (1975) has to be formulated more precisely than in the approach of BOX & TIAO (1975).

### 6. Illustrative analysis for an ARIMA(0,1,1)-model

The data set to illustrate the testing of intervention effects stems from a study by Meffert (see HOLTZMANN, 1963). One single schizophrenic patient's performance on a perceptual speed task was observed for 245 days (Fig. 4).

Fig. 4. Perceptual speed of a schizophrenic before, during and after drug administration.



During the first 60 days (phase I), the patient daily received a placebo drug, and during the second 60 days (phase II) chlorpromazine. From day 120 to day 180 (phase III) chlorpromazine was continued but with electroshock therapy superimposed. The final 65-day period (phase IV) was similar to the first. The analysis of the autocorrelations and partial correlations of every single phase shows that we can suggest an ARIMA(0,1,1)-model:

(6.1) 
$$(1-B)(Y_t - L) = (1 - \theta_1 B)a_t$$

The maximum likelihood (ML) estimation of  $\theta_1$  in phase I to III is approximately 0.76. Only for the last 65-day period  $\theta_1$  was estimated with 0.22.

The last estimation of  $\theta_1$  is significantly different from  $\hat{\theta}_1 = 0.76$ . This is an indication for a change in the model. Therefore we restrict the analysis to the effects of the first and second intervention.

The graph of the data shows some instability of level, which is typical for a nonstationary process. It seems also obvious that the treatments had markedly different effects upon the client. The introduction of the tranquilizer led to a downward shift in level of the series. The second intervention shows two effects, first a transient upward shift in level over a time period of 11 days and second a dynamic decreasing effect. Now we are testing the described intervention hypothesis with the model

(6.2) 
$$(1-B)(Y_t - L) = \omega_0 S_t^{(60)} + \omega_1 (S_t^{(120)} - S_t^{(131)}) + \frac{\omega_2}{1 - \delta_1 B} \cdot B^{12} \cdot S_t^{(120)} + (1 - \theta_1 B) a_t$$

Because of  $(1-B)(Y_t-L) = (1-B)Y_t$  we need not estimate the levelparameter L or the constant  $\theta_o$ . The ML estimation of the parameters are given in: MLE t-statistic

$$\begin{array}{c|ccccc} & & & & & & \\ \hat{\theta}_1 & & 0.764 & & 15.570 \\ & \hat{\omega}_0 & -21.986 & -3.656 \\ & \hat{\omega}_1 & & 9.137 & 2.040 \\ & \hat{\omega}_2 & -3.071 & -3.120 \\ & \hat{\delta}_1 & & 0.922 & 14.931 \end{array}$$

With the a priori information about  $\delta$  we can estimate the parameters L,  $\omega_0$ ,  $\omega_1$ ,  $\omega_2$  of the model (6.2) with the general linear model approach of GLASS et al. (1975). The designmatrix X is given by:

$$(6.4) \quad \underline{X} = \begin{bmatrix} 1 & \omega_{0} & \omega_{1} & \omega_{2} \\ 1 & 0 & 0 & 0 \\ 1 & 0 & 0 & 0 \\ \vdots & \vdots & \vdots & \vdots & \vdots \\ 60 & \\ 61 & \\ \vdots & \vdots & \vdots & \vdots & \vdots \\ 1 & 0 & 0 & 0 \\ \hline 1 & 1 & 0 & 0 \\ \hline 1 & 1 & 0 & 0 \\ \vdots & \vdots & \vdots & \vdots \\ 1 & 1 & 0 & 0 \\ \hline 1 & 1 & 0 & 0 \\ \hline 1 & 1 & 0 & 0 \\ \hline 1 & 1 & 0 & 0 \\ \hline 1 & 1 & 1 & 0 \\ \vdots & \vdots & \vdots & \vdots \\ 1 & 1 & 1 & 0 \\ 1 & 1 & 0 & 1 \\ 1 & 1 & 0 & 1 + \delta_{1} \\ \vdots & \vdots & \vdots & \vdots \\ 1 & 1 & 0 & \frac{57}{2} \delta_{1} \end{bmatrix}$$

Table (6.5) shows the results of the estimation with a priori known  $\delta_1 = 0.92$ :

- 1		MLE	t-statistic
	Ĺ	54.06	8.98
(6.5)	ŵο	-21.83	-3.63
	$\hat{\mathbf{\omega}}_1$	9.09	2.00
	$\hat{\omega}_2$	-3.09	-3.03

All the parameters are significantly different from zero on the  $\alpha = 0.05$  level.

#### Appendix A

The derivation of the  $\psi$ -weights:

The general autoregressive operator  $\varphi(B)$  is a polynominal in B of degree p+d:

A1) 
$$\begin{aligned} \phi(B) &= \phi_o - \phi_1 B - \phi_2 B^2 - \ldots - \phi_{p+d} B^{p+d} \\ \text{with} \qquad \phi_o &= 1 \end{aligned}$$

We have to derive the coefficients of this polynominal. Using the Binominal Expansion Theorem, we are able to express the difference operator as:

(A2) 
$$(1-B)^{d} = \binom{d}{0} - \binom{d}{1}B + \binom{d}{2}B^{2} - \binom{d}{3}B^{3} + \dots + (-1)^{d}\binom{d}{d}B^{d}$$
  
=  $\sum_{r=0}^{d} \binom{d}{r}(-1)^{r}B^{r}$ 

The autoregressive operator can be expanded as:

(A3) 
$$\Phi_{p}(B) = (1 - \Phi_{1}B - \Phi_{2}B^{2} - \dots - \Phi_{p}B^{p}) = \sum_{j=0}^{p} (-\Phi_{j})B^{j}$$
  
with  $-\Phi_{0} = 1$ 

We may combine (A2) with (A3), to get the general autoregressive operator

(A4)  $\varphi(B) = \left\{ \sum_{r=0}^{d} {d \choose r} (-1)^{r} B^{r} \right\} \cdot \left\{ \sum_{j=0}^{p} (-\Phi_{j}) B^{j} \right\}$ If we collect the coefficients of B<sup>k</sup> in (A4), we get

$$\varphi_{0} = -\Phi_{0} = 1$$

and for  $k = 1, \dots, p+d$ 

$$\begin{aligned} \varphi_{1} &= \varphi_{1} + \binom{d}{1} \\ \varphi_{2} &= \varphi_{2} - \binom{d}{1}\varphi_{1} - \binom{d}{2} \\ \text{(A5)} \qquad &\varphi_{3} &= \varphi_{3} - \binom{d}{1}\varphi_{2} + \binom{d}{2}\varphi_{1} + \binom{d}{3} \\ \varphi_{4} &= \varphi_{4} - \binom{d}{1}\varphi_{3} + \binom{d}{2}\varphi_{2} - \binom{d}{3}\varphi_{1} - \binom{d}{4} \\ &\vdots \\ \varphi_{k} &= \sum_{r=0}^{k} (-1)^{r+1}\binom{d}{r}(-\varphi_{k-r}) \qquad k = 0, 1, \dots, p+d \\ \text{where: } \binom{d}{r} &= 0 \text{ for } r > d \text{ and } -\varphi_{k-r} = 0 \text{ for } k-r > p \end{aligned}$$

Now we know the coefficients in  $\theta_q(B)$  and  $\varphi(B)$ . Using (3.19) we can collect the coefficients of  $B^k$  on the left and right side of (3.19) in a similar fashion as we did in (A1) – (A5). The coefficients of  $B^k$  can be arranged to a recursive equation-system:



Now the  $\psi$ -weights can be obtained recursively

As an example we want to derive the  $\psi$ -weights for the nonstationary ARIMA(1,1,1)-process

The coefficients of the general autoregressive operator are

(A9a) 
$$\begin{array}{l} \phi_{0} = 1 \\ \phi_{1} = (1 + \phi_{1}) \\ \phi_{2} = -\phi_{1} \end{array}$$

The same results could be obtained by (A5)

#### References

- AITKEN, A. C. (1935). On Least Squares and Linear Combination of Observations. Proceedings of the Royal Society of Edinburgh, 55, 42-48.
- BARLOW, D. H., and M. HERSEN (1973). Single-Case Experimental Designs, Archives of General Psychiatry, 29, 319–325.
- BOCK, R. D. (1975). Multivariate Statistical Methods in Behavioral Research. Mc Graw-Hill, New York.
- BOWER, C. P., PADIA, W. L., and G. V. GLASS (1974). TMS: Two FORTRAN IV Programs for the Analysis of Time-Series Experiments. Laboratory of Educational Research, Boulder, Co.
- Box, G. E. P., and G. M. JENKINS(1980). Time Series Analysis: Forecasting and Control. Holden-Day, San Francisco.
- BOX, G. E. P., and G. C. TIAO (1965). A Change in Level of a Nonstationary Time Series. Biometrika, 52, 181-192.
- Box, G. E. P., and G. C. TIAO (1975). Intervention Analysis with Applications of Economic and Environmental Problems, Journal of the American Statistical Association, 70, 70–79.
- CHASSAN, J. B. (1979<sup>2</sup>). Research Design in Clinical Psychology and Psychiatry. Halsted Press, New York.
- MCCLEARY, R., and R. A. HAY (1980). Applied Time Series Analysis For the Social Sciences. Sage Publications, London.
- FICHTER, M. M. (1979). Versuchsplanung experimenteller Einzelfalluntersuchungen in der Psychotherapieforschung. In: PETERMANN, F., and F. HEHL (ed.). Einzelfallanalyse, Urban & Schwarzenberg, Munich.
- GOLDBERGER, A. S. (1973). Structural Equation Models: An Overview. In: GOLDBERGER and DUNCAN (eds.). Structural Equation models in the Social Sciences, 1–18. Seminar Press, New York.
- GOTTMAN, J. M. (1981). Time-Series Analysis: A Comprehensive Introduction for Social Scientists. Cambridge University Press, Cambridge, Ma.
- GLASS, G. V., WILLSON, U. L., and J. M. GOTTMAN (1975). Design and Analysis of Time-Series Experiments. Colorado Associated University Press, Boulder, Co.
- HALL, R. V., FOX, R., WILLARD, D., GOLDSMITH, L., EMERSON, M., OWEN, M., DAVIS, F., and E. PORCIA (1971). The Teacher as Observer and Experimenter in the Modification of Disputing and Talking-out Behaviours. Journal of Applied Behaviour Analysis, 4, 141-149.
- HERSEN, M., and D. H. BARLOW (1976). Single-Case Experimental Designs: Strategies for Studying Behaviour Change. Pergamon Press, New York.

$$\begin{array}{l} (A9b) \quad \varphi_{1} = (-1)^{1} {\binom{d}{0}} (-\Phi_{1}) + (-1)^{2} {\binom{d}{1}} (-\Phi_{o}) = \Phi_{1} + 1 \\ \varphi_{2} = (-1)^{1} {\binom{d}{0}} (-\Phi_{2}) + (-1)^{2} {\binom{d}{1}} (-\Phi_{1}) + (-1)^{3} {\binom{d}{2}} (-\Phi_{o}) = -\Phi \\ = 0 \qquad \qquad = 0 \end{array}$$

Inserting the  $\phi_0$ ,  $\phi_1$ ,  $\phi_2$  into (A7) we get the  $\psi$ -weights

$\psi_o = 1$	$=\frac{1-\theta_1}{1-\varphi_1}\!+\!\frac{\theta_1-\varphi_1}{1-\varphi_1}\!\cdot\!\varphi_1^o$
$\psi_1 = \Phi_1 + (1 - \theta_1)$	$= \frac{1-\theta_l}{1-\varphi_l} \!+\! \frac{\theta_l-\varphi_l}{1-\varphi_l} \!\cdot\! \varphi_l^1$
$\psi_2 = \bigoplus_1 (\bigoplus_1 + (1 - \theta_1)) + (1 - \theta_1)$	$=\frac{1-\theta_1}{1-\varphi_1}\!+\!\frac{\theta_1-\varphi_1}{1-\varphi_1}\!\cdot\!\varphi_1^2$
$\psi_3 = \Phi_1(\Phi_1(\Phi_1 + (1 - \theta_1))) + (1 - \theta_1)$	$=\frac{1-\theta_1}{1-\varphi_1}\!+\!\frac{\theta_1-\varphi_1}{1-\varphi_1}\!\cdot\!\varphi_1^3$
1	
$\psi_j = \Phi_1^j + (1 - \theta_1) \sum_{i=0}^{j-1} \Phi_1^i$	$= \frac{1-\theta_1}{1-\varphi_1} + \frac{\theta_1-\varphi_1}{1-\varphi_1} \cdot \varphi_1^i$
(A10)	

HIBBS, D. A. JR. (1977). On Analyzing the Effects of Policy Intervention: BOX-JENKINS and BOX-TIAO vs. Structural Equation Models, in: HEISE, D. R. (ed.), Sociological Methodology, 137–179. Jossey Bass, New York.

- HIBBS, D. A. (1974). Problems of Statistical Estimation and Causal Inference in Time-Series Regression, in: COSTNER, H. S. (ed.), Sociological Methodology, 1973–1974. Jossey-Bass Publishers, San Francisco.
- HOLTZMANN, W. H. (1963). Statistical Models for the Study of Change in the Single Case, in: HARRIS, CH. (ed.), Problems in Measuring Change. The University of Wisconsin Press, Madison, Wisconsin.
- JENKINS, G. M. (1979). Practical Ecperiences with Modelling and Forecasting Time Series. GJP Ltd., Jersey, Channel Island.
- KAZDIN, A. E. (1976). Statistical Analysis for Single-Case Experimental Designs, in: HERSEN, M., and D. H. BARLOW (eds.), Single-Case Experimental Designs: Strategies for Studying Behaviour Change. Pergamon Press, New York.
- KEESER, W. (1979). Zeitreihenanalyse in der klinischen Psychologie: Ein empirischer Beitrag zur BOX-JENKINS Methodologie. Inaug. Diss. Munich.
- MÖBUS, C., and W. NAGL (1983). Messung, Analyse und Prognose von Veränderungen. In BREDENKAMP, J., and H. FEGER (ed.), Enzyklopädie der Psychologie, Bd. 5 "Hypothesenprüfung" in der Serie Forschungsmethoden, S. 239–470. Hogrefe, Göttingen.
- MOOSBRUGGER, (1978). Multivariate statistische Analyseverfahren. W. Kohlhammer, Stuttgart.
- REVENSTORF, D., and W. KEESER (1979). Zeitreihenanalyse von Therapieverläufen – ein Überblick. Urban & Schwarzenberg, Munich.
- TIMM, N. H. (1975). Multivariate Analysis With Applications in Education and Psychology. Brooks/Cole Publica., Monterey, Calif.
- TYLER, U. D., and G. D. BROWN (1968). Token Reinforcement of Academic Performance with Institutionalized Delinquent Boys. Journal of Educational Psychology, 59, 154-168.
- WOTTAWA, H. (1974). Das Allgemeine Lineare Modell Ein universelles Auswertungsverfahren. EDV in Medizin und Biologie, 3, 65-73.

Date of receipt: May 18th 1983.

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