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Application of GAs to Determine the Parameters of a Time Series Model

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ABSTRACT

We have witnessed in recent years a very rapid growth of work using genetic algorithms (GA). This trend can be observed in all areas of science economic. The aim of this article, we are interested in the application genetic algorithms for estimating the parameters of ARMA model (in the context of linear time series). To confirm the effectiveness of these new mechanisms, we apply both methods (the method of Box and Jenkins and her GA), to estimate the parameters of the model and compare the results obtained.

KEY WORDS - Genetic algorithm, parameters; ARMA models; Time series; method of Box and Jenkins.

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INTRODUCTION :-

The study of time series or time series corresponds to the statistical analysis of observations equally spaced in time. For the majority of these phenomena, there is often a dependence time between observations, which have autoregressive modeling: the past is used to explain the present and predict the future. Model and predict a time series is assumed in the most cases, to make assumptions about its behavior using as it is non-deterministic series he'll have the random component "varies, but not too much." This condition will result in the "stationary "Which implies a certain regularity of the process and allows deriving its asymptotic properties. When one has a series (X_t) in non-stationary stochastic, it should be modeled by a process $ARIMA(p; d; q)$, where d is the order of differentiation (or integration).

Having successfully transformed these data, the problem is to find a satisfactory $ARMA$ model and particularly as to determine p and q to find the autocorrelation functions and autocorrelation functions partial. The identification is mainly based on the analysis of the ACF (autocorrelation function) and $PACF$ (the partial autocorrelation functions) series considered.

The prediction method of *Box-Jenkins*^{1, 2} is particularly well suited to the treatment of series complex historical and other situations in which the Basic Law is not immediately apparent. However, as she deals with much more complicated situations, it is difficult to grasp the principles of this technique, as well as the limits of its application. In addition, the cost of me- Box-Jenkins method in a given situation is usually much higher than that of all other quantitative methods. We know that the applications of genetic algorithms are multiple: Medicine³, robotics⁴, analysis of time series^{5, 6}, image processing⁷

Our application examples helped us realize that the coding of data for model a problem is complex. On the other hand, we also saw difficulties effectively choose good parameter for the various operators (mutation, crossover, selection, replace placement). Choices in relation to the operators themselves are also manageable, knowing that some are most appropriate to the problem and let correlation means that they optimizer.

APPLICATION

We will study two sets of simulated and the other with real data by two methods-Box Jenkins' and 'Genetic Algorithm' and compare the results to make reliable forecasts.

Modeling the simulated series (B-J)

Begin with a series of simulated autoregressive process $AR(2)$, 300 observations. Table 1 gives the coefficients of the model^{8,9,10}.

Table 1 Coefficient model AR (2) of the series

Coefficient(s)	Estimate	Std. Error
ar1	1.5572	0.03665
ar2	-0.76809	0.03658

The graphic (1): Series (Y_t) and the series $(Y_t \text{ adjusted})$ adjusted by the model, shows the absence of a net difference between the two curves (we see a good fit).

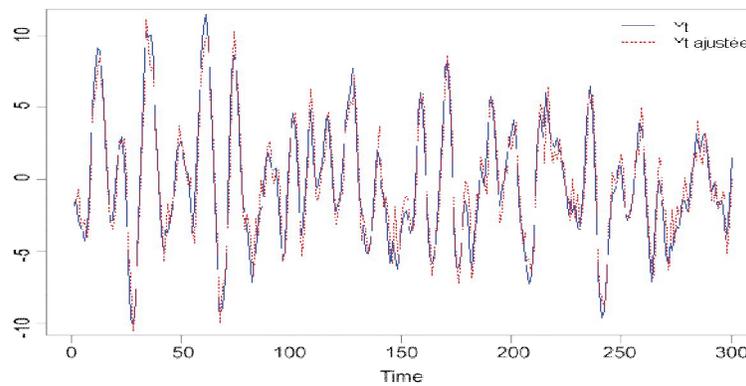


Figure 1 - Y_t and $Y_{t \text{ adjusted}}$

To validate the model we are going to study the residues (Table2).

Table 2 Tests on residue

Test	Formula	p -value
Kolmogorov-Smirnovles	lillie:test(residus)	0.08785
Student t	test(residus;mu = 0; conf:level = 0:95)	0.4671
Ljung - Box	Box.test(residus; lag = 50; type = "LjungBox")	0.1612

This allows us to conclude that the residues form a white noise, where model validation^{11,12,13}. Finally Y_t is adjusted by an $ARIMA(2;0;0)$ where $AR(2)$.

Modeling the simulated series (AG)

Now to the second method (Genetic Algorithm). it varies the number of observations, the size of the initial population (solutions ϕ_i) number of iteration of the algorithm genetic, crossover probability (from 1% to 100%) mutation probability (from 1% to 100%) and stop criterion (*mse* is minimum). (ϕ_1 and ϕ_2) Best, we got was (according to Table 3):

Table 3. The conditions for obtaining the best result (ϕ_1 and ϕ_2)

NBOBSERVATIONS = 298	number of observations
NBITERATIONS = 290	the initial population size
NBITERATIONS = 100	Iteration number of genetics algorithm
PROBCROISEMENT =90%	crossing probability
PROBMUTATION = 10%	probability of mutation
$\varepsilon = 0.000001$	stop criterion : $mse < \varepsilon$

$Best(\phi_1, \phi_2) = (1.5619365334, -0.7727730311)$, $mse = 1.6944448763$. The adjusted this model (GA) series is stationary over we see no difference between the two chronics (Figure 2).

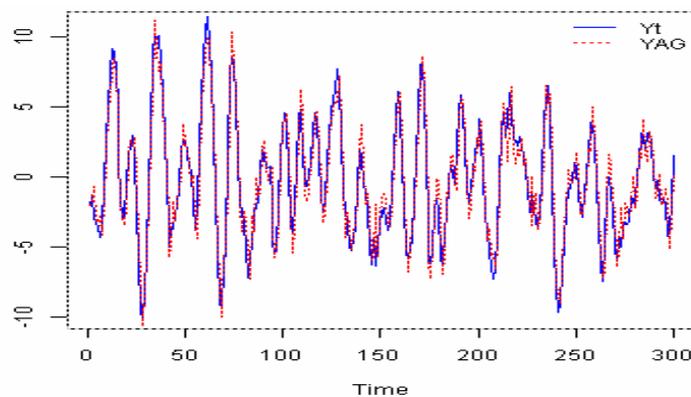


Figure 2 Yt and Yt ajus AG

Table 4 Tests on residue

Test	Formula	p -value
Kolmogorov-Smirnovles	lillie.test(residus)	0.3175
Student t	test(residus;mu = 0; conf:level = 0:95)	0.88
Ljung - Box	Box.test(residus; lag = 50; type = "LjungBox")	0.6273

This allows us to conclude that the residues form a white noise (Table 4), where model validation. We can have confidence in our predictions.

The graphic (3) Summarizes the quality of prediction and hence the performance of both models.

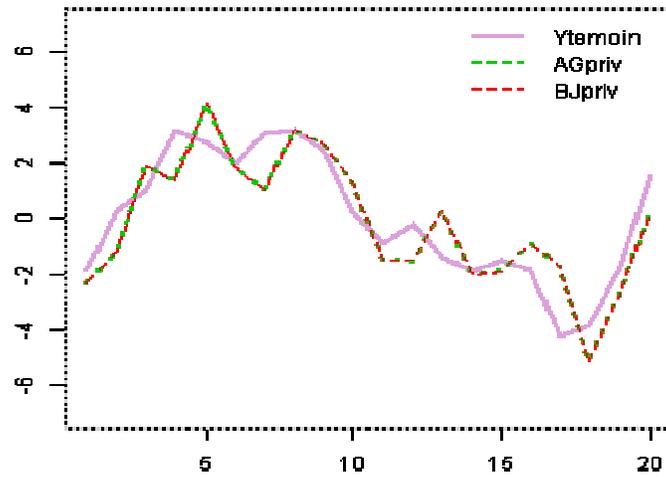


Figure 3 Performance of the two models GA & BJ

Visually, we see no difference between the two models. But the measures contained in the table(5), give preference to the model of Box and Jenkins.

Table 5 Conclusion comparison

	MAPE	MSE
B-J	1.158333	1.231707
GA	1.160062	1.233228

This time we discuss a real series, which represents the number of car accidents (weekly) for the years 1992 and 1995. We have 140 observations (source : Department of Statistics and O R ,Faculty of science, Kuwait University)¹⁴.

Modeling and Prediction of the real series (number of car accidents)

1.1 Method B- J

The graphic (4) Series, which represents the number of car accidents (weekly) for the years 1992 and 1995.

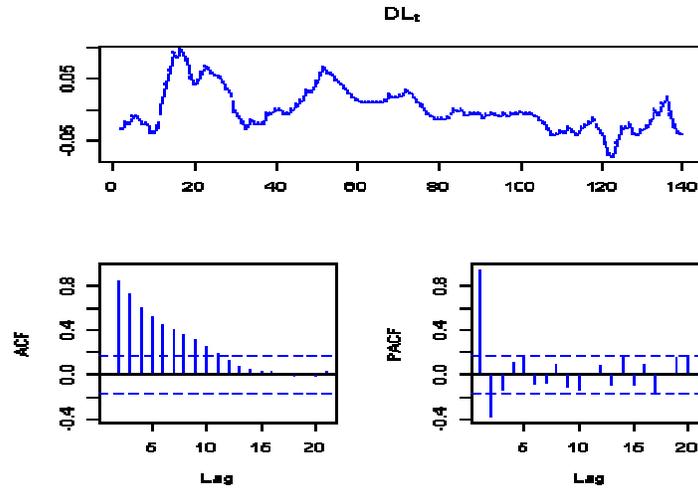


Figure 4 Y_t , ACF,PACF

After the two transformations : $L_t = \log(Y_t)$ and $DL = \text{diff}(L_t) = \text{diff}(\log(Y_t))$ we obtain a stationary series (DL) : p - value = 0:02066 test (adf.test). Best suited to this model is $ARIM(2;1;0)$. Table (6) gives the coefficients of the model¹⁴.

Table 6 Coefficient of the best model ARIMA (2; 1; 0) of the series

Coefficient(s)	Estimate	Std. Error
<i>ar1</i>	1.4139	0.0767
<i>ar2</i>	-0.4827	0.0772

Move on to the adjustment of our series by this model. According to the graphic (5), there is no differentiation between the two series.

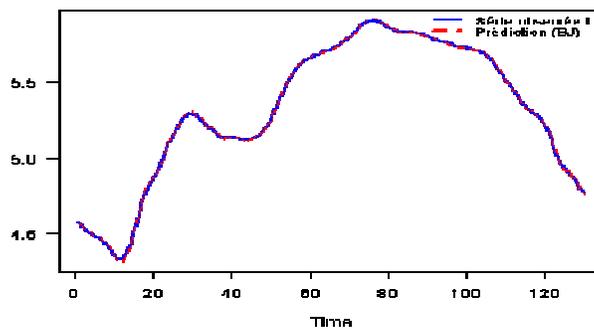


Figure 5 The adjustment of the series by this model (ARIMA (2; 1; 0))

To validate the model we will study the residues.

Table 7 Tests residues

Test	Formula	p -value
Kolmogorov-Smirnovles	lillie:test(residus)	0.1994
Student t	test(residus;mu = 0; conf:level = 0:95)	0.8935
Ljung - Box	Box.test(residus; lag = 50; type = "LjungBox")	0.9477

This allows us to conclude that the residues form a white noise (Table7), where model validation. We can have confidence in our predictions.

1.2 Method AG

Best $(\phi_1 \text{ and } \phi_2)$, Which was obtained under the conditions (8):

$$(\phi_1, \phi_2) = (1.3623294021, 0.4199999997)$$

Table 8 The conditions for obtaining the best result $(\phi_1 \text{ and } \phi_2)$

NBITERATIONS = 1000	the initial population size
NBITERATIONS = 100	nombre d'iteration of genetics algorithme
PROBCROISEMENT =90%	crossing probability
PROBMUTATION = 10%	probability of mutation
$\varepsilon = 0.000001$	stop criterion : $mse < \varepsilon$

The adjusted by the model and the real series are graphically almost identical graphic (6)

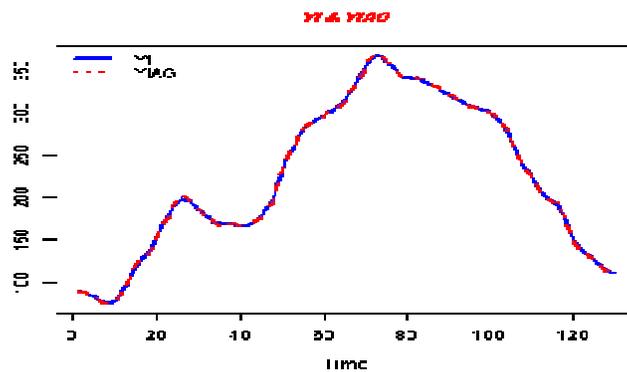


Figure 6 Yt and YtajusGA

To validate the model we will study the residues.

Table 9 Tests residues

Test	Formula	p -value
Kolmogorov-Smirnovles	lillie:test(residus)	0.3175
Student t	test(residus;mu = 0; conf:level = 0:95)	0.88
Ljung - Box	Box.test(residus; lag = 50; type = "LjungBox")	0.6273

This enables us to conclude that the residues form a white noise (Table 9), where model validation.

The graphic (7) summarizes the quality of prediction and hence the performance of both models.

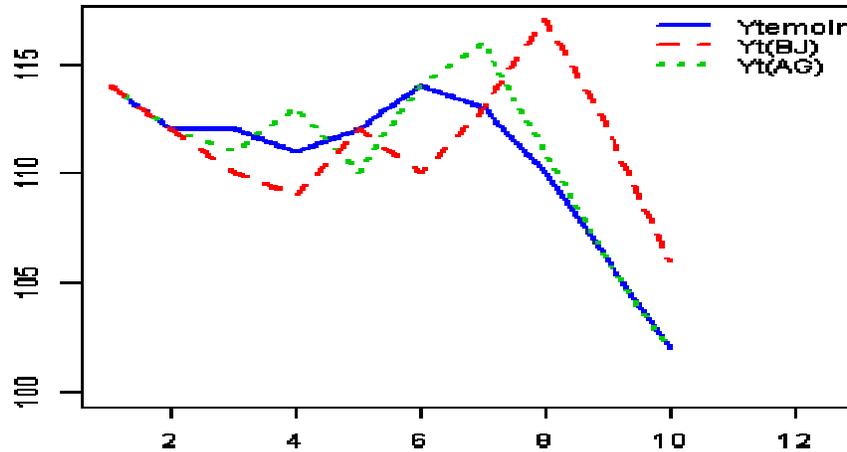


Figure 7 Performance of both models AG & BJ

Visually, there is a clear difference between the qualities of predictions, where the second validation method (GA).

Table 11 conclusion of comparison

	MAPE	MSE
B-J	1.158333	1.231707
GA	1.160062	1.233228

CONCLUSIONS

The first inspiration and driving force of this study were convinced that the sharing of experiences and knowledge between different fields of knowledge are an essential element of this enrichment even knowing . In the experimental part, we found that the use of GA Allow to obtain very good results in comparison with the results given by the method of Box Jenkins (in particular , for the actual chronic). Therefore, the conclusion of this study is not only proposal to continue the way of using Genetic Algorithms in the field of time series (parameter estimation), and thus predict, but we can say that genetic algorithms alone are not very effective in solving a problem. However,

they provide fairly quickly an acceptable solution. Nevertheless, it is possible to improve effectively enough by combining it with a deterministic algorithm.

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