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Environmental Problems

Екологічні проблеми

Volume 1 • Number 2



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MATHEMATICAL MODELLING OF THE POPULATION DYNAMICS OF HUNTING MAMMALS BASED ON RECURRENT EQUATION SYSTEM

I. Trach, V. Petruk, S. Kvaternyuk, T. Titov

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Abstract. Regression analysis of the population dynamics of hunting mammals based on Khmilnytskyi forestry was made. Correlation analysis of the observations results between the populations of spotted deer and wisent, spotted deer and wild boar, lepus and fox was made. Modelling of the population dynamics of hunting mammals was done using Mathcad based on the system of recurrent logistics equations and the adequacy of the model by the Pearson's criteria was proved.

Key words: mathematical modelling, population dynamics, hunting mammals, regression analysis, interspecies interaction, dynamics model.

Introduction. The use of mathematical modelling opens up new possibilities for creating a model of population dynamics and forecasting of the population, which greatly

facilitates the analysis of primary data. At the same time, there are some difficulties: a) simulation data requires a fairly long period of time (as the accuracy of the model is determined by the amount of data in number); b) it is difficult to reflect the impact of certain factors and interspecies relationships in conventional models [2–5].

1. The analysis of the populations of hunting mammals in Khmilnytskyi forestry

Based on the statistics of the population dynamics of hunting mammals in Khmilnitskyi forestry, a regression analysis was made using Statistica 6.1.Program. Exponential function was used as a regression model that corresponds to modelling the dynamics of individual populations without interspecies interaction. The results of the regression analysis are shown in Fig. 1.

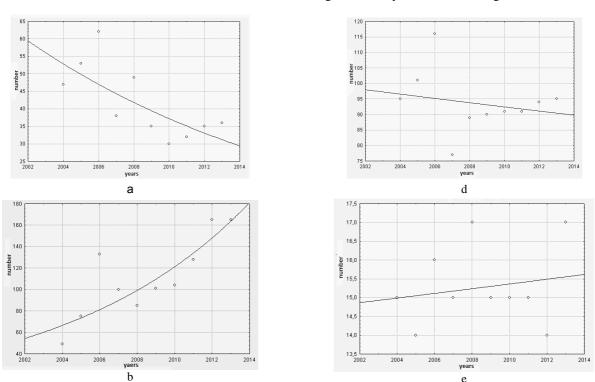
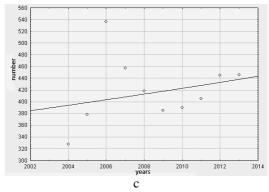
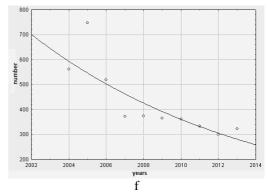


Fig. 1. Regression analysis of observations of population in Khmilnitskyi forestry: a – spotted deer, b – wild boar, c – European roe deer, d – wisent





Continuation Fig. 1. Regression analysis of observations of population in Khmilnitskyi forestry: e - fox, f - lepus.

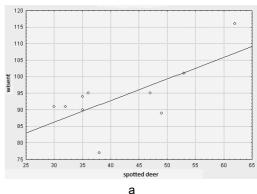
As a result of the regression analysis, mathematical models of changes in the number of individual populations of hunting animals in exponential approximation were obtained. Moreover, the function index includes parameters for fertility, mortality, migration and human impacts (hunting shooting, poaching, agricultural operations, transportation systems, etc.).

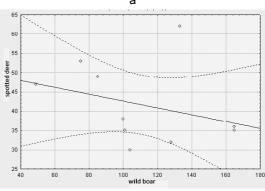
The correlation between the populations of certain species of hunting mammals in Khmilnitskyi forestry was analyzed. In particular, the dependence between the populations of spotted deer and wisent, fox and lepus, spotted deer and wild boar was set Fig. 2.

As a result of the correlation analysis between the populations of spotted deer and wisent in Khmilnitskyi forestry the correlation coefficient of 0.69 was obtained, which indicates close relationship between these populations. This is because the ecological niches of wisent and spotted deer coincide. Though, the population of spotted deer is much wider because of better adaptation to the existence in small forests. Spotted deer and wisent, in terms of trophic relationships, do not compete. In this case, such a high correlation coefficient is due to the fact, that wisent competes with deer, as a factor of disturbance. In spring and summer when ungulates are calving, deer are trying to avoid encounters with other animals. Herewith, in most cases overgrown cuttings and young forests are the places of the growth of the young ones. While eating, wisents use the same overgrown cuttings and young forests. The herd disperses throughout all the territory, making a loud noise and crackle. Other animals that brought the young ones try to leave this place, what is accompanied by the death of some young animals [1-5].

The lack of significant correlation relationship between the populations of fox and lepus in Khmilnitskyi forestry is explained by strong anthropogenic interference into ecological niches of animals and substantive hunting compared to interspecies interaction of "predator-prey". No evident correlation relationship between the populations of deer and wild boar was found, although their ecological niches coincide. This is because their number is small and they are not disturbing factors for each other. In terms of trophic relationships, they also do not significantly compete. Wild boar is able to live in any forestlands as well as

wetlands. It is omnivorous, which allows repeated changing of food base throughout the year.





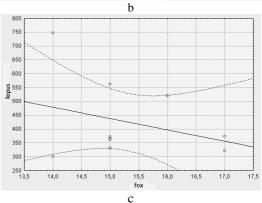


Fig. 2. Correlation analysis of observations between the populations in Khmelnytskiy forestry:

a – spotted deer and wisent, b – spotted deer and wild boar, c – lepus and fox

2. Modelling of the population dynamics of hunting mammals in Khmilnitskyi forestry using Mathcad

Mathematical model of dynamics of the amount of separate population under conditions with sufficient food, lack of overcrowding and enemies is described by the following equation:

$$N(t) = N_0 e^{r(t-t_0)};$$
 (1)

where N_0 – population amount at the initial time; r – specific rate of reproduction.

Equation (1) is received by solving Malthus differential equation $\frac{dN}{dt} = rN$.

Under adverse conditions, specific mortality rate d exceeds specific birth rate b, while the specific rate of reproduction r = d - b is negative.

When taking crowding into account, the population dynamics of a separate population is described by the following equation:

$$N(t) = \frac{K}{\ln\left(\frac{K - N_0}{N_0}\right) - r(t - t_0)};$$
(2)

where K – the maximum possible amount of the population.

Equation (2) is received by solving logistics differential equation $\frac{dN}{dt} = rN - \frac{r}{K}N^2$.

In consideration of interspecies interactions, it is necessary to solve a system of differential equations:

$$\begin{cases} \frac{dN_1}{dt} = r_1 N_1 - \frac{r_1}{K_1} N_1^2 + g_1 N_1 N_2; \\ \frac{dN_2}{dt} = r_2 N_2 - \frac{r_2}{K_2} N_2^2 + g_2 N_2 N_1. \end{cases}$$
(3)

where γ_1 and γ_2 – factors that take into account the interaction of species.

A more accurate study of the dynamics of interspecies interaction is possible with the use of systems

of nonlinear Lotka-Volterra differential equations, which take into account food digestion of the predator, predator's hunting strategy, defensive reaction of the victim, the presence of hiding places for the victim, etc.

With some loss of accuracy in the analysis and replacement of dt by Δt , the system of nonlinear differential equations may be replaced by the system of recurrent equations. This can significantly simplify the calculations and make them transparent and visible. For example, system (3) is converted into the following system:

$$\begin{cases} N_{i+1} = N_i + \left(r_n N_i - \frac{r_n}{K_n} N_i^2 + g_n N_i M_i \right); \\ M_{i+1} = M_i + \left(r_m M_i - \frac{r_m}{K_m} M_i^2 + g_m M_i N_i \right). \end{cases}$$
(4)

System (4) uses discrete time i which corresponds to the step of real time (average breeding season $T=min(T_n, T_m)$).

Modelling of the population dynamics in the ecosystem with six most common populations of hunting mammals was carried out. Initial data for the simulation are shown in Table 1.

Table 1
Initial data for the simulation
of the population dynamics

Species	Initial amount	Specific rate of change in population	Maximum population size	
Spotted deer	564	-0,0585	200	N1 _i
Wild boar	49	0,1003	160	N2 _i
European roe deer	384	0,0262	800	N3 _i
Wisent	96	-0,0073	100	N4 _i
Lepus	584	-0,0932	600	N5 _i
Fox	15	0,0041	17	N6 _i

Logistics system of recurrent equations for modelling the population dynamics:

$$\begin{bmatrix} N1_{i} + m1 \cdot N1_{i} - \frac{m1}{K1} \cdot \left(N1_{i}\right)^{2} + \gamma 12 \cdot N1_{i} \cdot N2_{i} + \gamma 13 \cdot N1_{i} \cdot N3_{i} + \gamma 14 \cdot N1_{i} \cdot N4_{i} + \gamma 15 \cdot N1_{i} \cdot N5_{i} + \gamma 16 \cdot N1_{i} \cdot N6_{i} + \gamma 17 \cdot N1_{i} \cdot N7_{i} \\ N2_{i+1} \\ N3_{i+1} \\ N3_{i+1} \\ N4_{i+1} \\ N5_{i+1} \\ N6_{i+1} \\ N7_{i+1} \end{bmatrix} = \begin{bmatrix} N1_{i} + m1 \cdot N1_{i} - \frac{m1}{K1} \cdot \left(N1_{i}\right)^{2} + \gamma 21 \cdot N1_{i} \cdot N2_{i} + \gamma 13 \cdot N1_{i} \cdot N3_{i} + \gamma 14 \cdot N1_{i} \cdot N4_{i} + \gamma 15 \cdot N1_{i} \cdot N5_{i} + \gamma 16 \cdot N1_{i} \cdot N6_{i} + \gamma 17 \cdot N1_{i} \cdot N7_{i} \\ N2_{i} + m2 \cdot N2_{i} - \frac{m2}{K2} \cdot \left(N2_{i}\right)^{2} + \gamma 21 \cdot N2_{i} \cdot N1_{i} + \gamma 22 \cdot N2_{i} \cdot N3_{i} + \gamma 24 \cdot N2_{i} \cdot N4_{i} + \gamma 25 \cdot N2_{i} \cdot N5_{i} + \gamma 26 \cdot N2_{i} \cdot N6_{i} + \gamma 27 \cdot N2_{i} \cdot N7_{i} \\ N3_{i} + m3 \cdot N3_{i} - \frac{m3}{K3} \cdot \left(N3_{i}\right)^{2} + \gamma 31 \cdot N3_{i} \cdot N1_{i} + \gamma 32 \cdot N3_{i} \cdot N2_{i} + \gamma 34 \cdot N3_{i} \cdot N4_{i} + \gamma 35 \cdot N3_{i} \cdot N5_{i} + \gamma 36 \cdot N3_{i} \cdot N6_{i} + \gamma 37 \cdot N3_{i} \cdot N7_{i} \\ N4_{i} + m4 \cdot N4_{i} - \frac{m4}{K4} \cdot \left(N4_{i}\right)^{2} + \gamma 41 \cdot N4_{i} \cdot N1_{i} + \gamma 42 \cdot N4_{i} \cdot N2_{i} + \gamma 43 \cdot N4_{i} \cdot N3_{i} + \gamma 45 \cdot N4_{i} \cdot N5_{i} + \gamma 46 \cdot N4_{i} \cdot N6_{i} + \gamma 47 \cdot N4_{i} \cdot N7_{i} \\ N5_{i} + m5 \cdot N5_{i} - \frac{m5}{K5} \cdot \left(N5_{i}\right)^{2} + \gamma 51 \cdot N5_{i} \cdot N1_{i} + \gamma 52 \cdot N5_{i} \cdot N2_{i} + \gamma 53 \cdot N5_{i} \cdot N3_{i} + \gamma 54 \cdot N5_{i} \cdot N4_{i} + \gamma 56 \cdot N5_{i} \cdot N6_{i} + \gamma 57 \cdot N5_{i} \cdot N7_{i} \\ N6_{i} + m6 \cdot N6_{i} - \frac{m6}{K6} \cdot \left(N6_{i}\right)^{2} + \gamma 61 \cdot N6_{i} \cdot N1_{i} + \gamma 62 \cdot N6_{i} \cdot N2_{i} + \gamma 63 \cdot N6_{i} \cdot N3_{i} + \gamma 64 \cdot N6_{i} \cdot N4_{i} + \gamma 65 \cdot N6_{i} \cdot N5_{i} + \gamma 67 \cdot N6_{i} \cdot N7_{i} \\ N7_{i} + m7 \cdot N7_{i} - \frac{m7}{K7} \cdot \left(N7_{i}\right)^{2} + \gamma 71 \cdot N7_{i} \cdot N1_{i} + \gamma 72 \cdot N7_{i} \cdot N2_{i} + \gamma 73 \cdot N7_{i} \cdot N3_{i} + \gamma 74 \cdot N7_{i} \cdot N4_{i} + \gamma 75 \cdot N7_{i} \cdot N5_{i} + \gamma 76 \cdot N7_{i} \cdot N6_{i} \end{bmatrix}$$

Simulation results are shown in Fig. 3.

Compare the results of modelling the dynamics of populations of spotted deer in Khmilnitskyi forestry from the system of recurrent equations with regression model based on the processing of observations using the exponential function. For comparison, we use the Pearson criteria. Since χ^2 =0,998, it indicates the adequacy of the model based on the system of recurrent equations to the results of observations (Fig. 4).

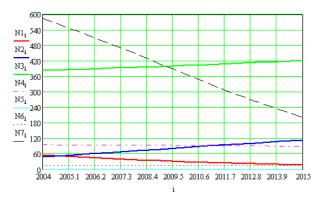


Fig. 3. The dynamics of the population amount of hunting mammals in Khmilnitskyi forestry

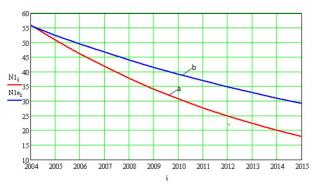
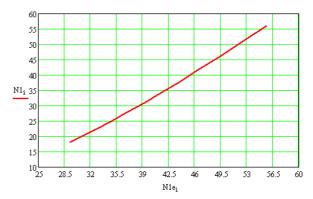


Fig. 4. Changes in the number of deer populations in Khmilnitskyi forestry (a – simulation results of equations (5), b) – exponential regression of observations)



The analysis of the adequacy of the data model of observations on the example of the dynamics of the populations of spotted deer (the Pearson criteria χ^2 =0,998).

4. Conclusions

The obtained model is somewhat arbitrary and cannot be used as a completely accurate prediction of the populations of hunting mammals. However, the scenarios of the population dynamics are recommended for the development of measures for the integrated management of the groups of hunting mammals both for Khmilnitskyi forestry hunting, and all hunting lands of Podillia region.

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