EFFECT OF EXOGENOUS GIVEN OXYTOCIN TO SOWS DURING PARITY,

ON THE PARTURITION TIME, STILLBIRTHS AND M.M.A. SYNDROM.
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In intensive modern swine production disturbances in physiological status of the organism which have direct or indirect influence on parturition are often encoutered. Some of these troubles are: prolonged time /more than 6 h/ of farrowing, increasing the stillbirth rate and a larger number /percentage ge/ of sows with mastitis, metritis agalac+ tiae syndrom.

Un the prevention of this poor-health status oxytocin parentterally given to sows is wi-

dely used.

syndrome.

Because of discrepant results obtained by differents authors, an attempt has been made to explain this problem.

Materials and methods

Observation were made in 3 industrial farms /A,B,C/ with 836 sows /primiparous and multiparous/ during their parturition period.
In farms A and B, animals were divided into 3 groups / I,II,III / and in farm C into 2 groups / I,II /. Groups I in all the farms were controls without any hormonal treatment. In groups II sows received 10 I.U. of synthetic oxytocin /Chemische Fabrik Gedeon Richter A.G. Budapest/ given intramuscullary /i.m./ after birth of the first piglet and eventually again, if interruption between succeding piglet delivery was longer than 1 hour. Sows from groups III /farms A, B/ received 2 ml of solution containing 10 I.U. oxytocin administered intranasally /i.n./ in the same manner as groups II. When no acceleration of delivery was observey ten to twenty minutes after the oxytocin applications, fetus position was examined and if necessary the fetus position was reposed. Farrowings of each group were observed and the following data were recorded: parturition time, stillbirths and incidence of M.M.A.

Results The results are presented in tables 1 and 2. Table 1. Influence of exogenous oxytocin on the parturition time.

Unit	Group	Nr of sows	lengh		of sows arturit 6-10	
A	I II III	100 100 100	8 34 12	75 57 66	15 9 12	0 0
В	III	100 100 100	41 45 47	40 47 44	14 8 9	5 0
C	I II III	118 118	32,6 39,7	53,4 51,8		0

Table 2. Influence of exogenous oxytocin, on stillbirth rate and the incidence of M.M.A. syndrome.

Group	Nr of sows	average litter size	Still Nr.	lbirths %	M.M.	
I	100	9,78	44	4,48	25	25
II	100	10,27	43	4,18	17	17
III	100	9,98	45	4,51	20	20

Continued tab 2.

II		10,77 9,41 10,17	68		
	118 118			4,98 4,23	

If we analize parturition times /tab 1/ we see that all sows which received oxytocin /i.m. of Risky farrowed shorter than 10 h; whilst sows in the control group farrowed longer than 10 h. In addition, the number of sows farrowed within the best possible time i.e. 1-6 h was by 5,5%-14% higher in the experimental groups than that in the con trols.

Similar results were obtained by /Tsoutsis et al, 1980, Zerobin 1972, Buning et al 1978, and Lee 1977/. It should be stressed that the above cited authors used different doses of oxytocin, from 0,1-0,2 I.U. administered i.v. by Zerobin to 40 I.U. administer red i.m. by Lee. Tsoutsis used 15-20 I.U. of oxytocin i.m.

The data contained in table 2 show that the stillbirth rate in the experimental sows from groups II was only about 0,30-0,97% lower than that in the control group. Sows in group III from farm 8 which received intranasal oxytocin had a lesser stillbirth rate only by 0, % in comparison to the control. Data in tab.2 regarding the results of M.M.A. syndrome are very interesting. As it has been shown in each experimental group much less cases of M.W.A. syndrome than in the control groups were observed. The differences in the activity of oxytocin

administered i.m. or i.n. were not essential, but it was always observed that oxytocin administered i.m. was slightly more effective.

Conclusions

- Oxytocin administered i.m. or i.n. in doses of 10 I.U. to sows during parturition shortens farrowing to less than 10 h, increases the number of sows farrowing in the optimal physiological time 1-6 h, and reduces the occureance of M.M.A. syndrome. - Oxytocin administered to sows as above does not lesser the rate of stillbirth.

Selected references: Buning J., Duckhert J.: Mh. Vet-med 1978, 33,446. Curtis S.L.: J. Anim. Sci. 1974, 38,1031. Duensing R.: Ph.). Tier-Sci. 1974, 28, 1031. Duensing R.: Ph. D.Tier-arztl.Hochschule Hannover 1970.Lee Co.: Korea J.Vet.Res. 1977, 17, 12. Randall G.C.B. Biol. Neonate 1979, 36, 55. Tsoutsis J.J. Proc. Congr. I.P.V.S. Commanagen 1980. Lrathall a.s.: Vet.Rec. 1971, 89, 61. Zerobin K., Spörri R.: Adv. vet.sci.comp.med. 1972, 16, 521, Zerobin E.: Proc. Congr. I.R. I.P.V.S. 1980.