Animal Science Papers and Reports vol. 22 (2004) no. 1, 57-64 Institute of Genetics and Animal Breeding, Jastrzebiec, Poland

Growth rate and carcass quality in relation to *GH/MspI* and *GH/Hae*II PCR-RFLP polymorhism in pigs*

Mariusz Pierzchała, Tadeusz Blicharski, Jolanta Kurył

Polish Academy of Sciences Institute of Genetics and Animal Breeding, Jastrzebiec, 05-552 Wólka Kosowska, Poland

(Received January 9, 2004; accepted February 27, 2004)

The objective of this study was to investigate the effect of two single nucleotide polymorphisms in growth hormone (GH) gene on performance traits in pigs. The animals (n= 320) belonged to four groups of commercial crosses being the progeny of crossbred Polish Landrace × Polish Large White sows and Polish Landrace, Polish Large White, Duroc or Pietrain boars. Genotypes of growth hormone gene (GH) were established with PCR-RFLP technique using *MspI* and *HaeII* endonucleases. The *GH/MspI* and *GH/HaeII* genotypes were found significantly related to the weight of ham, weight of ham meat and ham content of carcass. Moreover, nearly significant differences between *GH/MspI* genotypes were found for mean fat thickness (from five measurements), fat thickness at lower back (point K2) and over the loin, and for loin eye height. The results presented allow to assume that near the analysed mutation the QTLs for performance traits in pigs are located.

KEY WORDS: carcass / gene polymorphism / growth hormone / pig

The "candidate gene" approach is purposeful when a gene is known to function in such a way that it may explain genetic variation in traits of interest. It is well known that growth hormone (GH) is one of the most important factors for growth and development of animal cells. Porcine GH is synthesized as 190-amino acid peptide and secreted pulsatile by acidophilic or somatotropic cells of the anterior pituitary [Kato *et al.* 1990]. The GH gene (GH) is assigned to pig chromosome 12 [Thomsen *et al.* 1990, Yerle *et al.* 1993, Chowdhary *et al.* 1994]. GH has long been known to have a significant beneficial effect on carcass quality in pigs. Treating pigs with exogenous porcine GH improves daily live weight gain, feed efficiency and lean content of carcass [Chung

^{*}Supported by the State Committee for Scientific Research, grant No. 6 PO6D 020 25.

and Etherton 1986, Etherton et al. 1986]. Comparison between pig lines selected for live weight gain and backfat thickness showed correlation of both high growth rate and low backfat thickness with higher blood GH level [Rindberg Lund-Larsen and Bakke 1975, Althen and Gerrits 1976]. Because of physiological importance of GH the GH gene was the objective of several studies as the candidate gene for performance traits in farm animals [Zwierzchowski et al. 2002, Ge et al. 2003]. The effect of GH variants on performance traits in pigs have been reported, among others, by Casas-Carrillo et al. 1997, Knorr et al. 1997, Krenkowa et al. 1999, Pierzchała et al. 1999, Cheng et al. 2000, and Putnova et al. 2001].

The goal of the pig industry is to produce the high quality, lean pork. The most economically important traits are sow fertility, growth rate, feed conversion, meat content of carcass, as well as meat quality. Growth process and cell proliferation in animals are regulated by multitude of physiological pathways among which the somatotropic axis plays a key role. Thus the differences in frequency of restriction sites identified as GH gene polymorphisms may explain growth rate and carcass composition differences occurring between pigs.

In this study we tried to evaluate the effect of the porcine GH/MspI and GH/HaeII polymorphisms on growth rate and carcass quality in commercial pig crosses.

Material and methods

The pigs belonged to four commercial lines obtained by crossing of purebred boars (Polish Landrace, Polish Large White, Duroc, Pietrain) with crossbred (Polish Landrace × Polish Large White) sows. Eighteen growth rate and carcass quality traits were measured in a total of 320 animals.

Polymorphism of growth hormone gene (GH) was identified with PCR-RFLP using primers and procedure of Kirkpatrick [1992]: MspI within second intron (A-284 and 222 bp, B - 222, 147 and 137 bp) and *Hae*II within the second exon (A - 506 bp, B - 173 and 333 bp). Considered was also the effect of *RYR1* genotype.

Statistical analysis using the least squares (SAS Institute Inc., Cary, USA) was performed to find significant differences between GH genotypes in 18 recorded traits of performance. The following model was used:

$$Y_{ijklm} = \mu + G_i + B_j + S(B)_{jk} + R_l + (GB)_{ij} + (GR)_{il} + \beta(wc_{ijklm} - wc) + e_{ijklm}$$

where: Y_{ijklm} - trait measured on *ijklm*-th animal;

 μ – overall mean:

- G_i = effect of *i*-th genotype (*GH/Msp*I = 1,2,3; *GH/Hae*II = 1,2,3);
- B_j = effect of *i*-th breed of a sire (j = 1, 2, 3, 4);

 $S(B)_{jk}$ - nested effect of *j*-th sire within the *k*-th breed;

 R_l = effect of *l*-th RYR1 genotype (l = 1, 2, 3);

 $(GB)_{ij}$ - effect of interaction *i-th* genotype × *j*-th breed;

 $(GR)_{il}$ - effect of interaction *i*-th genotype × *l*-th RYR1 genotype;

 β^{-} regression coefficient on cold carcass weight;

 e_{ijklm} - random effect.

Results and discussion

AA, AB, and BB frequencies of *GH/MspI* were found 3.8, 28.4 and 67.8%, and of *GH/Hae*II – 22.2, 50.6, and 27.2%, respectively (figures not tabulated).

Within *GH/MspI locus* (Tab. 1) higher values for loin eye height, weight of ham, weight of ham meat, and ham content of carcass were found significantly (P<0.05) associated with genotype BB, while higher mean fat thickness (from five measurements) and backfat thickness at point K2 nearly significantly (P<0.1) with AA genotype.

Within *GH/Hae*II *locus* (Tab. 2) higher values of weight of ham and weight of ham meat (P<0.05) as well as ham content of carcass (P<0.01) were found associated with genotype AA. There were no differences for fatness related to *GH/Hae*II genotypes.

QTL mapping performed by Korwin-Kossakowska *et al.* [2001] within the Polish Pig Genome Mapping Project provides information on localization of genes affecting fat deposition (abdominal fat) on chromosome 12 near the microsatellite markers S0083 and S0090. These microsatellites are linked to *GH* gene [Larsen *et al.* 1995, Rohrer *et al.* 1994, 1996, Archibald *et al.* 1994, Korwin-Kossakowska *et al.* 1999]. Studies performed by Arbona *et al.* [1988] demonstrated that pigs selected for increased growth rate had higher basal plasma GH concentration than unselected pigs, whereas in a study by Saleri *et al.* [2001] growth rate in pigs was significantly correlated not with GH, but with insuline-like growth factor I (IGF-I) and insuline-like growth rate and selected carcass traits in pigs.

Associations of another two *GH* variants with pig performance traits were analysed by Knorr *et al.* [1997] in two resource families – Wild boar × Pietrain (W × P) and Meishan × Pietrain (M × P). Significant associations between eight carcass fatness traits and *GH/HinP1*I and *GH/Apa*I variants were found only in family M × P. This corroborates the results of Larsen *et al.* [1995] who also could not find such relations in W × P pigs. In our earlier study on the reference family Zlotnicka Spotted × Polish Large White [Pierzchala *et al.* 1999], as well as in that performed by Krenkova *et al.* [1999] similar relations between *GH/Hae*II genotype and fat and meat deposition traits were found. Moreover, Křenkova *et al.* [1999] showed the unfavourable effect of both AA *GH/Msp*I and AA *GH/Hae*II genotypes on mean fatness and leaness of carcass, while

60								
			C.M.	M			raca ku resteriPi	6.
Trac					ź			
	ğ	B	ğ	Ð	ğ	Ð	aav v	aa-av
Control on sometheous, by	Ş	1	ě,	~	Ę	-		
Puctorations (Tront) reservation (Inc)	2	E o	116	- Laa	161	30	- 0	
Bei Redidnem (cn)	2	E		D D D		3		
Puthetinan utkow had – 51 (cn)	21	R	ΪĻ	80	4	100		
Purchaterea advocrised = 51 (cn)	196	A a	2		Ē	001		ā
Puththan utwarked = 51 (cm)	21	h o	μ	Πą	161	0 0		
Puththan evolution (cn)	-	H o	\$	80	141			
Puthting exceledation)	17 -	R 0		80	115	100		
Herteoffange (or)	U 9	А О	979 979		661	80		
Then of lange (or)	9 61) 0	ş	Πg	911	80		
At scatter ferren (1)		р —	L 1	016	T T T	1-0		
At scored have (16)	n A		ЯR Д	6-0 0	ЦЯЦ	ā		
Sau altanın cıram (19	21	ī	9	рП	\$ [11 0	- 0	
The effect have we have been a [2]	A R	þ	ĝ	121	11M	2	9 0	2
The age of the second of the (g)	۹L	e	2 <u>9</u>	쿿	3	ę		
The effect have (p)	-81	ß		ЦĢ	1 M I	<u>\$</u>	-	8
Trepto Navi kora (p	Ħ	-		≝	¢[]]	≏		
Tegeoften noch)		Ħ	Ī	E		\$	ġ	

Table I. Leasingare now (LSA) un ter nucled ann (S2) fir pofimare and una date G/MAM grappenn

60

				3			đ	iraca ka	6
Trac							Ĩ	a na su la	
	8	B	ļ	E	1	B	ann	A MEB	98 - 94
buly ive voeteeun je)	1		Ĭ	-0	ŝ	o			
National Trady recurrent (24)	цтг	B o	Ë	50	Ę	9 0			
ket fied dinem (cn)	116	8 0	116	50	1	9 0			
Networken science but - 61 (cn)		= 0	11LI		176	10			
hetetran edoverbed =61 (cn)	9	0 0	2		III	ΠQ			
tetrations advice but = E1 (cn)	Ĕ	1 0	31	Πđ	161	0 16			
hethethen everlen (cn)	\$ -	d o	12 1	\$	12				
hetetra extraction (a)	2	= 0		80	3	10			
tegteoflancy: (ov)	-2	1 0	6 24	Πđ		11 0			
Pichi d'Isinge (or)	f ¢		976	Ē	ŝ	11 0			
de se concerce d'e aroun (19)	115	р 0	21%	20	412				
descenario (16)	9 2	d 0	TB CT	0 0	П£	ŝ			
Part of American (19		н о	51 T L	ЪЦ	22	à	20	80	
Stepted Nan vertex basel (2)	100-1	2	TRIV	2	92 2	Ē		2 0	
Pt gte d'harveware fie (e)	ULI	F	9	5	-121	9			
Pterstearthern (p)	97LI	9		Ħ		â		80	
StyteChen kena (g	Ē	3			11-6 1	Ä	20		
Ptetechen work)	9	<u>9</u>	201	Ξ	-15	Ä		9 0	

i na Maru a I
CV NAME AL
r pathmu
2(8)5
न्त्रमा त्म
urd dar -
0veril en
Ā

in the study of Pierzchała *et al.* [1999] the respective unfavourable effect was found of only AA *GH/Hae*II, the AA *GH/MspI loci* affecting both traits positively. Relation between carcass quality and both *GH/MspI* and *GH/Hae*II *loci* is evident also from the present results (Tab. 1 and 2). Significant and positive effect of BB *GH/MspI* genotype was found on ham weight, and especially on ham meat weight, as well as nearly significant on ham content of carcass, and backfat thickness. Positive effects on ham weight, ham meat weight, and ham content of carcass were also found of AA *GH/Hae*II genotype.

As far as *GH/MspI locus* is concerned, the present results partially corroborate those reported by Křenkova *et al.* [1999], but are opposite to our earlier data [Pierzchała *et al.* 1999]. Considering *GH/Hae*II *locus*, differences between genotypes point out to trends opposite to those obtained by us earlier, as well as presented recently by Kurył *et al.* [2003]. Importance of this region of genome for carcass performance traits in pigs was also mentioned by Geldermann *et al.* [2003] where several QTLs were localized on chromosome 12, and especially near the *GH* gene [Yue *et al.* 2003].

The results obtained in several studies on the relationship between *GH* gene variants and carcass traits, sometimes slightly opposite to each other cannot, so far, be definitely summarized, but there is enough evidence that the linkage of analysed *GH* gene variants with other mutation (causal) is highly probable.

REFERENCES

- ALTHEN T.G., GERRITS R.J. 1976 Pituitary and serum growth hormone levels in Duroc and Yorkshire swine genetically selected for high and low backfat. *Journal of Animal Science* 42, 1490-1497.
- ARBONA J.R., MARPLE D.N., RUSSEL R.W., RAHE C.H., MULVANEY D.R., SARTIN J.L. 1988 – Secretory patterns and metabolic clearence rate of porcine growth hormone in swine selected for growth. *Journal of Animal Science* 66, 3068-3072.
- 3. ARCHIBALD A.L., HALEY C.S., BROWN J.F., COUPERWHITE S., MCQUEEN H.A., NI-CHOLSON D., COPPIETERS W., VAN DE WEGHE A., STRATIL A., WINTERO A.K., FRED-HOLM M., LARSEN N.J., NIELSEN V.H., MILAN D., WOLOSZYN N., ROBIC A., DALENS M., RIQUET J., GELLIN J., CARITEZ J.-C., BURGAUD G., OLLIVIER L., BIDANEL J.-P., VAIMAN M., RENARD C., GELDERMANN H., DAVOLI R., RUYTER D., VERSTEGE E.J.M., GROENEN M.A.M., DAVIES W., HOYHEIM B., KEISERUD A., ANDERSSON L., ELLEGREN H., JOHANSSON M., MARKLUND L., MILLER J.R., ANDERSON DEAR D.V., SIGNER E., JEFFREYS A.J., MORAN C., LE TISSIER P., MULADNO, ROTSCHILD M.F., TUGGLE C.K., VASKE D., HELM J., LIU H.-C., RAHMAN A., YU T.P., LARSON R.G., SCHMITZ C.B., 1995 The PigMap consortium linkage map of the pig. *Mammalian Genome* 6, 157-175.
- CASAS-CARRILLO E., PRILL-ADAMS A., PRICE S.G., CLUTTER A.C., KIRKPATRICK B.W., 1997 – Relationship of growth hormone and insulin-like growth factor-1 genotypes with growth and carcass traits in swine. *Animal Genetics* 28, 88-93.
- CHOWDHARY B.P., THOMSEN P.D., HARBITZ I., LANDSET M., GUSTAVSSON I., 1994

 Precise localization of the genes for glucose phosphate isomerase (GPI), calcium release channel (CRC), hormone-sensitive lipase (LIPE), and growth hormone (GH) in pigs, using nonradioactive in situ hybridization. *Cytogenetics and Cell Genetics* 67, 211-4.
- CHENG W.T., LEE C.H., HUNG C.M., CHANG T. J., CHEN C.M. 2000 Growth hormone gene polymorphisms and growth performance traits in Duroc, Landrace and Tao-Yuan pigs. *Theriogenology* 54, 1225-1237.

- CHUNG C.S., ETHERTON T.D., 1986 Stimulation of swine growth by porcine growth hormone. Journal of Animal Science 60, 118
- GE W., DAVIS M.E., HINES H.C., IRVIN K.M., SIMMEN R.C., 2003 Association of single nucleotide polymorphisms in the growth hormone and growth hormone receptor genes with blood serum insulin-like growth factor I concentration and growth traits in Angus cattle. *Journal of Animal Science* 81, 641-8.
- GELDERMANN H., MÜLLER E., MOSER G., REINER G., BARTENSCHLAGER H., CEPICA S., STRATIL A., KURYL J., MORAN C., DAVOLI R., BRUNSCH C., 2003 – Genome wide linkage QTL mapping in porcine F2 families generated from Pietrain, Meishan and Wild Boar crosses. *Journal of Animal Breeding and Genetics* 120, 363-393.
- ETHERTON T.D., WIGGINS J.P., CHUNG C.S., EVOCK C.M., REBHUN J.F., WALTON P.E., 1986 – Stimulation of pig growth performance by porcine growth hormone and growth releasing factor. *Journal of Animal Science* 63, 1389-1399
- KATO Y., SHIMOKAWA N., KATO T., HIRAI T., YOSHIHAMA K., KAWAI H., HATTORI M., EZASHI T., SHIMOGORI Y., WAKABAYASHI K., 1990 – Porcine growth hormone: molecular cloning of cDNA and expression in bacterial and mammalian cells. *Biochimica et Biophysica Acta* 1049, 290-293
- 12. KNORR C., MOSER G., MÜLLER E., GELDERMANN H., 1997 Associations of GH gene variants with performance traits in F2 generation. *Animal Genetics* 28, 124-128.
- KIRKPATRICK B.W., 1992 HaeII and MspI polymorphisms are detected in the second intron of the porcine growth hormone gene *Animal Genetics* 23, 180-181.
- 14. KORWIN-KOSSAKOWSKA A., PIERZCHAŁA M., CYMEROWSKA-PROKOPCZYK I., SZYDŁOWSKI M., KURYŁ J., ŻURKOWSKI M., KAMYCZEK M., JANIK A. 2001 – The Polish "Pig Genome Mapping" project. XIII. Identification of quantitative trait loci affecting carcass fat deposition. *Animal Science Papers and Reports* 19, 27-42.
- KORWIN-KOSSAKOWSKA A., PIERZCHAŁA M., KURYŁ J., ZWIERZCHOWSKI L., CYME-ROWSKA-PROKOPCZYK I.., SIADKOWSKA E., 1999 – Polymorphism of porcine growth hormone gene and its linkage to microsatellites S0083 and S0090. *Journal of Applied Genetics* 40, 85-91.
- KŘENKOVA L., KUCIEL L., URBAN T., 1999 Association of the RYR1, GH, LEP and TF genes with carcass and meat quality traits in pigs. *Czech Journal of Animal Science* 44, 481-486
- KURYŁ J., KAPELAŃSKI W., PIERZCZHAŁA M., BOCIAN M., GRAJEWSKA S., 2003 A relationship between genotypes at *GH* and *LEP loci* and carcass meat and fat deposition in pigs. *Animal Science Papers and Reports* 21, 15-26.
- LARSEN N.J., ELLEGREN H., BRAUNER NIELSEN P., ANDERSSON L., 1995 Genetic variation at growth hormone locus in wild pig intercross; test of association to phenotypic traits and linkage to the blood group D locus. *Theoretical and Applied Genetics* 91, 1047-1077.
- PIERZCHAŁA M., KORWIN-KOSSAKOWSKA A., ZWIERZCHOWSKI L., LUKASZEWICZ M., ZIEBA G., KURYL J., 1999 – *Hae*II and *MspI* polymorphism of growth hormone gene in pigs and its association with production traits. *Czech Journal of Animal Science* 44, 441-445.
- PUTNOVÁ L., KŘENKOVÁ L., VRTKOVÁ I., DVOŘÁK J., PIETRUSZKA A., CZARNECKI R., 2001– Association od the DdeI growth hormone gene polymorphism with some performance traits in Polish Large White and Czech Large White × Polish Large White pigs. *Journal of Applied Genetics* 42, 317-324
- RINGBERG LUND-LARSEN T., BAKKE H., 1975 Growth hormone and somatomedin activities in lines of pigs selected for rate of gain and thickness of backfat. *Acta Agriculturae Scandinavica* 25, 231-234.
- 22. ROHRER G.A., ALEXANDER L.J., HU Z., SMITH K.P.L., KEELE J.W., BEATTIE C.W., 1996 A comprehensive map of the porcine genome *Genome Research* 6, 371-391.

- 23. ROHRER G.A., ALEXANDER L.J., KEELE J.W., SMITH T.P.L., BEATTIE C.W., 1994 A microsatellite linkage map of the porcine genome. *Genetics* 136, 231-245.
- 24. SALERI R., BARATTA M., MAINARDI G.L., RENAVILLE R., GIUSTINA A., QUINTAVALLA F., TAMANINI C., 2001 IGF-I, IGFBP-2 and 3 but not GH concentrations are different in normal and poor growing piglets. *Reproduction, Nutrition, Development* 41, 163-172.
- THOMSEN P.D., FREDHOLM M., CHRISTENSEN K., SCHWERIN M., 1990 Assignment of the porcine growth hormone gene to chromosome 12. *Cytogenetics and Cell Genetics* 54, 92-4.
- 26. YERLE M., LAHBIB-MANSAIS Y., THOMSEN PD., GELLIN J., 1993 Localization of the porcine growth hormone gene to chromosome 12p1.2→p1.5. *Animal Genetics* 24, 129-31.
- YUE G., SCHRÖFFEL JR J., MOSER G., BARTENSCHLAGER H., REINER G., GELDERMANN H., 2003 – Linkage and QTL mapping for *Sus scrofa* chromosome 12 *Journal of Animal Breeding and Genetics* 120, 95-102.
- ZWIERZCHOWSKI L., KRZYZEWSKI J., STRZAŁKOWSKA N., SIADKOWSKA E., RYNIE-WICZ Z., 2002 – Effects of polymorphism of growth hormone (GH), Pit-1, and leptin (LEP) genes, cow's age, lactation stage and somatic cell count on milk yield and composition of Polish Black-and White cows. *Animal Science Papers and Reports* 20, 213-227.

Mariusz Pierzchała, Tadeusz Blicharski, Jolanta Kurył

Tempo wzrostu i cechy jakości tuszy świń w powiązaniu z restrykcyjnym polimorfizmem PCR-RFLP genu hormonu wzrostu – *GH/MspI* i *GH/ Hae*II

Streszczenie

Celem badań było określenie zależności między cechami użytkowymi świń a polimorfizmem genu hormonu wzrostu (*GH*). Materiał badawczy stanowiły zwierzęta pochodzące z krzyżowania knurów czystych ras (pbz, wbp, Duroc, Pietrain) z lochami mieszańcowymi (wbp \times pbz). Analizą objęto 18 cech tempa wzrostu i jakości tuszy 320 zwierząt.

Istotne zależności między analizowanymi genotypami *GH/Msp*I i *GH/Hae*II odnotowano dla masy szynki, masy mięsa szynki oraz udziału masy szynki w tuszy (%). Ponadto stwierdzono bliskie istotności zależności między genotypem *GH/Msp*I a średnią grubością słoniny z pięciu pomiarów, grubością słoniny mierzoną na krzyżu w punkcie K2, grubością słoniny okrywającej polędwicę oraz wysokością oka polędwicy. Prezentowane wyniki wraz z wcześniejszymi doniesieniami potwierdzają tezę, że w rejonie chromosomu 12 obejmującym *locus GH* znajdują się QTLs, które istotnie wpływają na poziom cech użytkowych świń.