Comparison of Preservation of Liquid Boar Semen between Lactose-Egg Yolk and Bütschwiler Diluents

Park, C. S., Y. M. Cheon and Z. Xu*

Department of Animal Science, College of Agriculture, Chungnam National University, Taejon 305-764, Korea

돼지 액상정액 보존을 위한 Lactose-Egg Yolk와 Bütschwiler 희석액의 비교

朴昌植・千庸敏・徐 直*

忠南大學校 農科大學 畜產學科

요 약

본 연구는 돼지 액상정액을 인공수정용 100ml 플라스틱 병에 보존하면서 Lactose-Egg yolk 희석액과 Bütschwiler 희석액 간에 보존 온도별 차이를 조사하고, Lactose-Egg yolk 희석액에서의 글리세롤 농도의 효과를 조사하여 돼지 액상정액을 좀더 장기간 사용할 수 있는 방법을 찾고자 실시하였다. 돼지 액상정액을 5℃ 냉장고에 보존하면서 조사한 바에 의하면, 37℃에서 0.5 및 2시간 배양 후의 정자 운동성은 전체 보존기간동안 Lactose-Egg yolk 희석액이 Bütschwiler 희석액보다 유의하게 (P<0.05) 높게 나타났고, 정상첨체비율은 두 희석액간에 차이가 없었다. 돼지 액상정액을 15℃에 보존하면서 조사한 바에 의하면, 3일부터 7일 보존시까지 정자 운동성과 정상첨체비율에 있어서 Bütschwiler 희석액이 Lactose-Egg yolk 희석액보다 유의하데 높게 나타났다. Lactose-Egg yolk 희석액을 이용한 돼지 액상정액의 글리세롤 농도의 효과에 있어서는 최종 글리세롤 농도가 0, 1, 3 및 5% 보다 2%일 때 가장 높은 정자 운동성과 정상첨체비율을 나타내었다. 분만율, 복당 생존자돈수 그리고 출생시 평균 생시체중은 Lactose-Egg yolk 희석액과 Bütschwiler 희석액간에 차이가 없었다. 이상의 연구 결과를 종합해 볼 때 Lactose-Egg yolk 희석액은 5℃에서 Bütschwiler 희석액은 15℃에서 6~7일 동안 돼지 액상정액을 보존할 수 있었다.

I. INTRODUCTION

Several diluents for liquid boar semen are used throughout the world by commercial artificial insemination centers. Among them, Modena (Moretti, 1981) is the most widely used in Korea. Summermatter (1984) modified Modena diluent by adding cysteine and BSA, and increasing the proportion of glucose, and used it as Bütschwiler. The above diluents have generally been the most successful within the storage

이 논문은 한국과학재단의 "1994년도 한국과학자 중국 파견연구"의 연구결과임.

^{*} Tian Jin Institute of Animal Husbandry and Veterinary Science, Tian Jin 300112, China.

temperature of 15 to 18°C. Chung et al. (1989), using BF_5 diluent for pellet freezing, found that liquid semen can be stored at 5°C for 9 days in 5 ml straw. However, the processing method of liquid semen in 5 ml straw was not simple for practical purpose.

Lactose-Egg yolk diluent is used for the freezing diluent of straws. Commercial use of Lactose-Egg yolk diluent is likely to continue until significantly better freezing diluents are forthcoming.

The objectives of this study were to find out methods of long-term use of liquid semen in 100 ml plastic bottle for artificial insemination and to investigate differences between Lactose-Egg yolk and Bütschwiler diluents according to storage temperature, and effect of final glycerol concentration in Lactose-Egg yolk diluent.

II. MATERIALS AND METHODS

1. Semen collection

Semen was collected from three Large White boars twice weekly. The filtered sperm-rich fraction was collected by the gloved-hand technique into a 250 ml insulated vacuum bottle. The sperm-rich fractions of ejaculates with greater than 85% motile sperm and NAR acrosome were used in this experiments.

2. Semen evaluation

Sperm motility and NAR acrosome of ejaculated semen and diluted liquid semen were evaluated by microscopical examination. The sperm concentration was estimated with a hemocytometer. One ml semen was pipetted into test tube containing 4 ml of Beltsville Thawing Solution (BTS). An 1 ml of the aliquot of the diluted semen was then added to 2 ml of 1% glutaraldehyde in BTS for acrosome morphology evaluation. The remaining 4 ml of diluted semen was

incubated at 37° C. After 0.5 and 2h incubation, 6 μ aliquots were transferred onto glass slides and 18×18 mm cover-slips were applied.

The percentage of motile sperm was estimated at 37°C by phase-contrast microscope at 250×. The acrosome morphology of 100 sperm per sample at 0.5 and 2h after incubation at 37°C was evaluated by phase contrast microscopy at 1,000×. Acrosomes were differentially categorized into four morphological classes: normal apical ridge(NAR), damaged apical ridge (DAR), missing apical ridge(MAR) and loose acrosomal cap(LAC) as described by Pursel et al.(1972) and Pursel and Park(1987).

3. Semen processing

The compositions of Lactose-Egg yolk, Bütschwiler and BTS diluents are shown in Table 1.

1) Processing method of liquid semen preserved at 5%

- (1) Semen is slowly cooled to room temperature by 2h after collection.
- (2) Plastic bottle of 100ml for artificial insemination(Mini tube, Germany) is filled with 10ml of semen (4~5×10⁸ sperm/ml) at room temperature.
- (3) Lactose-Egg yolk or Bütschwiler diluent (first diluent) of 10ml are added in the plastic bottle with 10ml of semen at room temperature.
- (4) The plastic bottle with the first diluted semen is cooled in a refrigerator to 5°C over a 2h period and Lactose-Egg yolk or Bütschwiler + 6% glycerol diluents(second diluent) of 10ml are added to the first diluted semen.
- (5) The diluted semen of 30ml in the plastic bottle is stored in the refrigerator until inseminated. The diluted semen is assessed in the laboratory for motility and acro-

Table 1. The composition of diluents

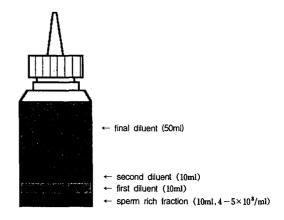
Ingredient	Lactose-Egg yolk ¹	Bütschwiler ¹	BTS ³
Glucose		3.5(g)	3.7(g)
Lactose hydrate	11.0(g)		
Sodium citrate		0.69	0.6
Sodium bicarbonate		0.10	0.125
TES			
EDTA, disodium		0.235	0.125
KCl			0.075
Hepes		0.95	
Egg yolk(ml)	25.0		
Tris, buffer		0.565	
Citric acid		0.315	
Cysteine		0.0054	
BSA		0.3	
Penicillin	0.01	0.01	0.01
Streptomycin	0.01	0.01	0.01
Distilled water(ml)	100	100	100

Lactose-Egg yolk or Bütschwiler + 6% glycerol²

some using the method described the above.

- (6) At insemination, the plastic bottle with 30ml of the diluted semen is rediluted with 50ml BTS diluent.
- (7) The dilution method of liquid semen preserved at 5°C is shown in Fig. 1.

- (1) Semen is slowly cooled to room temperature by 2h after collection.
- (2) Plastic bottle of 100 ml for artificial insemination is filled with 10 ml of semen $(4\sim5\times10^8~{\rm sperm/ml})$ at room temperature.
- (3) Lactose-Egg yolk or Bütschwiler diluents (first diluent) of 20 ml are added in the plastic bottle with 10 ml of semen at room



temperature.

(4) The diluted semen of 30 ml in the plastic bottle is stored at 15°C until inseminated.

¹ Lactose-Egg yolk and Bütschwiler = first diluent

² Lactose-Egg yolk or Bütschwiler + 6% glycerol = second diluent

³ BTS = final diluent.

The diluted semen is assessed in the laboratory for motility and acrosome using the methods described the above.

(5) At insemination, the plastic bottle with 30ml of the diluted semen is rediluted with 50ml BTS diluent.

4. Artificial insemination

Estrus was checked twice daily at 09:00 and 15:00h. The first insemination was done with liquid boar semen stored for 6 days at 22~26h after detection of standing estrus, using the spiral-tip catheter(Melrose and O'Hagan, 1961). The second insemination was done with the liquid boar semen stored for 7 days at 12h after the first insemination.

III. RESULTS AND DISCUSSION

1. Sperm viability in Lactose-Egg yolk or Bütschwiler diluents at 5°C

The percentages of motility on the liquid boar semen diluted with Lactose-Egg yolk or Bütschwiler diluents and preserved at 5°C are presented in Table 2-1.

The motility after 0.5 and 2h incubation at 37°C, respectively, was significantly higher for sperm preserved in Lactose-Egg yolk diluent

than for sperm preserved in Bütschwiler diluent from 1 to 7 day of storage. The sperm motility in Lactose-Egg yolk diluent steadily declined from 83.1% and 81.0% at 1 day of storage to 73. 1% and 65.0% after 7 day of storage at 0.5 and 2h incubation, respectively.

The percentages of NAR acrosome on the liquid boar semen diluted with Lactose-Egg yolk or Bütschwiler diluent and preserved at 5°C are presented in Table 2-2. The NAR acrosome in Lactose-Egg volk diluent steadily declined from 83.4% and 71.0% at 1 day of storage to 52.9% and 49.8% after 7 day of storage at 0.5 and 2h incubation, respectively. The NAR acrosome in Lactose-Egg yolk diluent after 0.5 and 2h incubation at 37°C, respectively, during preservation periods was similar to that in Bütschwiler diluent. The above results were similar to the report of Chung et al. (1989) using BF₅ diluent in 5ml maxi-straw at 5°C. So the diluted sperm with Lactose-Egg yolk diluent in 100ml plastic bottle will be stored for 7 day at 5°C. However, the sperm with Bütschwiler diluent was not used at 5°C preservation temperature.

2. Sperm viability in Lactose-Egg yolk or Bütschwiler diluents at 15°C

The percentages of motility on the liquid boar

Diluent	Incubation		Storage length	of semen (day)1	
	time(h)	1	3	5	7
Lactose-Egg yolk	k 0.5	83.1 ± 1.53 ^a	80.0 ± 3.16 a	78.5 ± 1.80 a	73.1 ± 1.83^{a}
Bütschwiler		35.0 ± 5.47 b	23.1 \pm 1.60 $^{\rm b}$	22.6 ± 6.04 b	18.1 ± 3.65 b
Lactose-Egg yolk	0	81.0 ± 3.66 a	76.3 ± 1.75 a	75.9 ± 1.57 a	65.0 ± 3.16 a
Bütschwiler	2	31.3 ± 5.96 b	22.4 ± 1.86 b	21.5 ± 7.20 ^b	15.0 ± 2.14 b

¹ Values were observed after 0.5 and 2h incubation in water bath of 37℃. Mean±S.E. for 6 ejaculates from Large White boar.

a. b Means in the same column with different superscripts at 0.5 and 2h incubation time differ significantly (P < 0.05), respectively.

Table 2-2. Comparison of percentage of NAR on the liquid boar semen diluted with two diluents and preserved at 5°

D3 4	Incubation	Storage length of liquid semen (day)1				
Diluent	time(h)	1	3	5	7	
Lactose-Egg yolk	٥۶	83.4 ± 6.58	72.0 ± 4.52	63.1 ± 2.31	52.9 ± 2.46	
Bütschwiler	0.5	84.1 ± 2.63	77.0 ± 5.17	68.0 ± 6.34	57.3 ± 6.37	
Lactose-Egg yolk	2	71.0 ± 6.27	64.0 ± 4.62	56.3 ± 3.02	49.8 ± 2.26	
Bütschwiler	2	75.3 ± 1.96	66.1 ± 5.06	59.2 ± 3.81	51.3 ± 5.12	

¹ Values were observed after 0.5 and 2h incubation in water bath of 37°C. Mean ± S.E. for 6 ejaculates from Large White boar.

semen diluted with Lactose-Egg yolk or Bütschwiler diluents and preserved at 15° C are preserved at 15° C are preserved in Table 3-1.

The motility after 0.5 and 2h incubation at 37°C, respectively, was significantly higher for sperm preserved in Bütschwiler diluent than for sperm preserved in Lactose-Egg yolk diluent from 3 day of storage. The sperm motility in Bütschwiler diluent steadily declined from 85.0% and 81.8% at 1 day of storage to 66.8% and 58.6% after 7 day of storage at 0.5 and 2h incubation, respectively.

The percentages of NAR acrosome on the liquid boar semen diluted with Lactose-Egg yolk or Bütschwiler diluents and preserved at 15°C are presented in Table 3-2. The NAR acrosome after 0.5 and 2h incubation at 37°C, respectively, was

significantly higher for sperm preserved in Bütschwiler diluent than for sperm preserved in Lactose-Egg yolk diluent from 3 day of storage. The NAR acrosome in Bütschwiler diluent steadily declined from 89.0% and 88.1% at 1 day of storage to 63.1% and 58.8% after 7 day of storage at 0.5 and 2h incubation, respectively.

Weitze(1991) reported that the percentages of motility and NAR acrosome after storage for 144h at 15°C were $75.0 \sim 80.8\%$ and $87.6 \sim 89.6\%$ in BW₂₅ diluent, and $73.3 \sim 81.7\%$ and $87.9 \sim 90.3\%$ in Androhep diluent, respectively. The BW₂₅, Androhep and Reading(Revell and Glossop, 1989) diluents showed high motility and NAR acrosome as compared with the Bütschwiler diluent. In future, we need to compare the above diluents under the same environment. In this

Table 3-1. Comparison of percentage of motility on the liquid boar semen diluted with two diluents and preserved at $15\,^{\circ}$

Diluent	Incubation	Storage length of liquid semen (day) ¹				
	time(h)	1	3	5	7	
Lactose-Egg yolk	0.5	85.0 ± 3.16 a	$71.8 \pm 1.83^{\ b}$	53.6 ± 4.84 ^b	20.0 ± 3.16 b	
Bütschwiler	0.5	85.0 ± 3.16 ^a	80.0 ± 3.16 a	75.0 ± 3.16 ^a	66.8 ± 4.83 a	
Lactose-Egg yolk	2	80.0 ± 6.32 a	63.6 ± 3.66 b	34.5 ± 3.20 b	10.0 ± 7.74 b	
Bütschwiler	۷	81.8 ± 1.83 a	73.6 ± 1.86 a	65.0 ± 3.16 a	58.6 ± 3.66 a	

¹ Values were observed after 0.5 and 2h incubation in water bath of 37℃. Mean±S.E. for 6 ejaculates from Large White boar.

a. b Means in the same column with different superscripts at 0.5 and 2h incubation time differ significantly (P< 0.05), respectively.</p>

Table 3-2. Comparison of percentage of NAR on the liquid boar semen diluted with two diluents and preserved at $15\,^{\circ}$

Diluent	Incubation	Storage length of liquid semen (day)				
	time(h)	1	3	5	7	
Lactose-Egg yolk	0.5	86.1 ± 1.47^{a}	69.0 ± 1.26^{b}	$23.5 \pm 5.64^{\text{b}}$	8.0 ± 2.89^{b}	
Bütschwiler		89.0 ± 1.67^{a}	84.1 ± 2.04^{a}	71.8 ± 4.57^{a}	63.1 ± 1.94^{a}	
Lactose-Egg yolk	9	84.1 ± 3.25^{a}	$56.8 \pm 3.65^{\text{b}}$	$15.8 \pm 3.71^{\rm b}$	3.6 ± 3.14^{b}	
Bütschwiler	2	88.1 ± 1.83^{a}	73.8 ± 2.99^{a}	65.1 ± 4.11^{a}	58.8 ± 1.60^{a}	

¹ Values were observed after 0.5 and 2h incubation in water bath of 37°C. Mean±S.E. for 6 ejaculates from Large White boar.

study, the diluted sperm with Bütschwiler diluent in 100 ml plastic bottle will be stored for 7 days at 15°C. However, the sperm with Lactose-Egg yolk diluent was not used after 3 days at 15°C preservation temperature.

Effect of glycerol concentration of liquid semen in Lactose-Egg yolk diluent

The percentages of motility and NAR acrosome on the liquid boar semen diluted different

glycerol concentration in Lactose-Egg yolk diluent are presented in Table 4-1 and 4-2.

As shown in Table 4-1, the motility after 0.5 and 2h incubation at 37°C, respectively, was significantly higher for sperm preserved in 0, 1 and 2% final glycerol concentrations than for sperm preserved in 3 and 5% final glycerol concentrations.

As shown in Table 4-2, NAR acrosome after 0. 5 and 2h incubation at 37°C, respectively, was

Table 4-1. Comparison of the glycerol concentration on the percentage of motility of liquid boar semen diluted with Lactose-Egg yolk diluent

Final glycerol concentration (%)	Incubation	Stora	ge length of liqu	id semen at 5 $^\circ{ m C}$ (day)¹
	time (h)	1	3	5	7
0		80.0 ± 3.16^{a}	63.1 ± 48.3^{a}	63.1 ± 1.83^{a}	48.1 ± 4.83^{a}
1		81.3 ± 3.66^{a}	68.1 ± 1.83^{a}	65.0 ± 3.16^{a}	53.1 ± 7.96^{a}
2	0.5	80.0 ± 3.16^{a}	68.1 ± 1.83^{a}	66.3 ± 1.86^{a}	55.0 ± 3.36^{a}
3		73.1 ± 4.83^{ab}	$56.3 \pm 1.86^{\circ}$	55.0 ± 3.16 ^b	26.3 ± 3.66 ^b
5		68.1 ± 7.96 ^b	$51.3 \pm 1.86^{\circ}$	$46.3 \pm 1.86^{\circ}$	$18.1 \pm 4.83^{\text{b}}$
0		71.3 ± 1.86^{ab}	53.1 ± 1.83^{a}	46.3±4.84°	30.0 ± 6.32^{a}
1		76.3 ± 3.66^{a}	56.3 ± 3.66^{a}	51.3 ± 6.59^{a}	31.3 ± 4.59^{a}
2	2	75.0 ± 0.00^{a}	61.3 ± 3.66^{a}	55.6 ± 5.73^{a}	35.0 ± 3.47^{a}
3		66.3 ± 1.86 ^{bc}	36.3 ± 1.86 ^b	28.1 ± 4.83^{b}	15.0±3.16 ^b
5		$58.1 \pm 1.83^{\circ}$	$31.8 \pm 3.65^{\text{b}}$	25.0 ± 3.16 ^b	13.1 ± 1.83^{b}

¹ Values were observed after 0.5 and 2h incubation in water bath of 37℃. Mean±S,E, for 6 ejaculates from Large White boar.

a, b Means in the same column with different superscripts at 0.5 and 2h incubation time differ significantly (P<0.05), respectively.</p>

a. b. c Means in the same column with different superscripts at 0.5 and 2h incubation time differ significantly(p<0.05), respectively.</p>

Table 4-2. Comparison of percentage of NAR on the glycerol concentration of liquid boar semen diluted with Lactose diluent

Final glycerol	Incubation	Stora	ge length of liquid semen at 5℃ (day)¹		
concentration (%)	time (h)	1	3	5	7
0		81.1 ± 1.94 ^{bc}	$67.5 \pm 4.46^{\text{b}}$	57.6±2.25 ^b	48.0±2.19 ^b
1		$82.8 \pm 2.63^{\mathrm{ab}}$	74.8 ± 1.60^{a}	69.0 ± 1.67^{a}	58.8 ± 2.78^{a}
2	0.5	87.0 ± 3.16^a	$78.0 \pm 2.75^{\circ}$	68.0 ± 3.94^{a}	58.9 ± 1.63^{a}
3		83.1 ± 3.65^{ab}	71.3 ± 3.01^{ab}	$62.1 \pm 2.04^{\mathrm{ab}}$	45.0 ± 2.89^{b}
5		$77.3 \pm 1.03^{\circ}$	$60.1 \pm 2.63^{\circ}$	$46.3 \pm 1.03^{\circ}$	$26.0 \pm 1.89^{\circ}$
0		82.0 ± 1.09^a	62.0±1.26 ^b	44.1 ± 2.22°	36.1 ± 3.18^{a}
1		84.0 ± 3.94^{a}	71.0 ± 3.16^a	61.0 ± 2.19^{a}	42.8 ± 2.85^{a}
2	2	86.1 ± 2.63^{a}	70.8 ± 2.99^{a}	62.1 ± 3.12^{a}	44.0 ± 2.19^{a}
3		75.0 ± 2.19 ^b	57.3 ± 2.58 ^{bc}	54.0 ± 2.19^{b}	36.1 ± 5.74^{a}
5		$52.1 \pm 1.60^{\circ}$	$52.1 \pm 1.60^{\circ}$	$41.3 \pm 1.36^{\circ}$	$16.0 \pm 2.89^{\text{b}}$

¹ Values were observed after 0.5 and 2h incubation in water bath of 37°C. Mean±S,E, for 6 ejaculates from Large White boar.

significantly higher for sperm preserved 1 and 2% final glycerol concentrations than for sperm preserved in 0, 3 and 5% final glycerol concentrations.

Glycerol concentration affects post-thaw survival of sperm frozen by either the straw or pellet method. The general view is that relatively low concentrations of glycerol(1 to 3%) are required for maximum post-thaw survival(Pursel et al., 1978; Paquignon, 1985; Almlid and Johnson, 1988). Graham and Crabo(1972), comparing post-thaw viablity of boar sperm frozen with 0, 1, 2, or 5% glycerol, found that both motility and NAR acrosome of sperm decreased at 5%

level. In the present study, values for motility were maximum at 1 or 2% glycerol and for NAR acrosome were maximum at 2% glycerol.

4. Fertilizing capacity of Lactose-Egg yolk and Bütschwiler diluents

Farrowing rate, litter size and average pig weight at birth for semen diluted in Lactose-Egg yolk or Bütschwiler diluents are presented in Table 5.

Farrowing rate, litter size and average pig weight at birth did not differ significantly between Lactose-Egg yolk and Bütschwiler diluents. The above results were similar to the report

Table 5. Fertilizing capacity of liquid semen stored in 100ml plastic bottle with Lactose-Egg yolk of 2% glycerol or Bütschwiler diluents

Diluent	No. of gilts	Farr	owed	No. of pigs born	Average pig weight
	inseminated1	No.	%	alive per litter	at birth(kg)
Lactose-Egg yolk	16	14	87.5	10.4 ± 1.88	1.32 ± 0.49
Bütschwiler	16	13	81.3	10.6 ± 1.96	1.34 ± 0.36

¹ Gilts were inseminated twice with liquid boar semen stored at 5℃ in Lactose-Egg yolk or 15℃ in Bütschwiler diluents for 6~7 days. Sperm concentration was 50×10⁸ sperm /80ml in 100ml plastic bottle.

a, b, c Means in the same column with different superscripts at 0.5 and 2h incubation time differ significantly(p<0.05), respectively.</p>

of Park et al. (1992) on fertilizing capacity of liquid boar semen depending on sperm concentration in 5ml maxi-straw. In this study, we found out that liquid boar semen can be stored for $6\sim7$ days at 5° c in Lactose-Egg yolk diluent and 15° c in Bütschwiler diluent.

IV. SUMMARY

This study was done to find out the methods of long-term use of liquid boar semen in 100 ml plastic bottle for artificial insemination and to investigate differences between Lactose-Egg yolk and Bütschwiler diluents according to storage temperature, and effect of final glycerol concentration in Lactose-Egg yolk diluent, Liquid boar semen diluted with Lactose-Egg yolk diluent showed significantly higher sperm motility (p<0.05) after 0.5 and 2h incubation at 37°C than Bütschwiler diluent at all storage length when it was preserved in the 5°C refrigerator. The NAR acrosome in Lactose-Egg volk diluent after 0.5 and 2h incubation at 37°C, respectively, during preservation periods was similar to that in Bütschwiler diluent. When liquid boar semen was preserved at 15°C, liquid boar semen in the Bütschwiler diluent showed significantly higher percentages of sperm motility and NAR acrosome from third day to seventh than that in Lactose-Egg yolk diluent. In the effect of final glycerol concentration of liquid boar semen in the Lactose-Egg yolk diluent, the final glycerol concentration of 2% showed higer percentages of sperm motility and NAR acrosome than that of 0, 1, 3, and 5%. Farrowing rate, litter size and average pig weight at birth did not differ significantly between Lactose-Egg yolk and Bütschwiler diluents. As a result of this study, we found out that liquid boar semen can be stored for 6~7 days at 5°C in Lactose-Egg yolk diluent and at 15°C in Bütschwiler diluent,

V. REFERENCES

- 1. Almlid, T. and L. A. Johnson. 1988. Effects of glycerol concentration, equilibration time and temperature of glycerol addition on post-thaw viability of boar spermatozoa frozen in straws. J. Anim. Sci., 66: 2899-2905.
- Chung, H. K., H. K. Kim, M. S. Ko, I. C. Kim, J. S. Choi, K. W. Lee, D. S. Son, H. Kim, S. H. Chee and C. S. Park. 1989. Studies on fertilizing capacity and survival of liquid boar semen in 5ml maxi straws. Korean J. Anim, Sci., 31(3): 158-161.
- Graham, E. F. and B. G. Crabo. 1972. Some factors influencing the freezing of boar spermatozoa. 7th Int. Congr. Anim. Repord. and A. I., Munich, pp 1629-1632.
- Melrose, D. R. and C. O'Hagan. 1961. Investigation into the techniques of insemination in the pig. Proc. IVth Int. Congr. Anim. Report. The Hague, IV, 855.
- Moretti, 1981. Cited by Johnson. L.A. and I. G. Aalbers (1984). Artificial insemination of swine: fertility using several liquid semen diluents. In proceedings of the 8th International Pig Verterinary Society Congress, Ghent, Belgium, 293.
- Paquignon, M. 1985. Freezing and thawing extenders of boar spermatozoa. In: Johnson, L.A. and K. Larsson. (Ed.) Deep Freezing of Boar Semen. Swedish Univ. of Agric. Sci., Uppsala, Sweden. pp 129-146.
- Park, C. S., S. W. Han, J. S. Soh, D. I. Kim, H. K. Chung and C. G. Ryu. 1992. Study on fertilizing capacity of liquid boar semen depending on sperm concentration in 5ml maxi-straw. Korean J. Anim. Sci., 34(2) :97-100.
- 8. Pursel, V. G., L. A. Johnson and G. B. Rampacek. 1972. Acrosome morphology of boar

- spermatozoa incubated before cold shock. J. Anim, Sci. 34: 278-283.
- 9. Pursel, V. G. and C. S. Park. 1987. Duration of thawing on post thaw acrosome morphology and motility of boar spermatozoa frozen in 5 ml maxi-straw. Theriogenology, 28: 683-690.
- Pursel, V. G., L. L. Schulman and L. A. Johnson. 1978. Effect of glycerol concentration on frozen boar sperm. Theriogenology, 9: 305-312.
- 11. Revell, S. G. and C. E. Glossop, 1989, A

- long-time ambient temperature diluent for boar semen. Anim. Prod., 48: 579-584.
- Summermatter, P. 1984. Möglichkeiten zur Verbesserung der Konservierbarkeitävon Ebersperma. Verhandlungsber. Verterinär-Humanmedizinische Gemeinschaftstagung, Hannover, FRG, 72-75.
- Weitze, K. F. 1991. Long-term storage of extended boar semen. In: Johnson, L.A and D. Rath. (Ed.) Boar Semen Preservation II. Paul Pariey Scientific Publishers, Berlin and Hamburg. Germany. pp 231-253.