Summer trial, UCR

Isobac 20, a product of Nationwide Chemical Corporation, was reported to give good control of damping-off as an infurrow spray, or as a pre-emergence spray at ground cracking or during the post-emergence period. The active ingredient of Isobac 20 is monosodium salt of 2.2 methylenebis (3.4.6-trichlorophenol). Isobac was compared with Panogen 15 2 oz, Panogen plus Vitavax 75W 8 oz, Ceresan L 2 oz plus Demosan 65W 10 oz, Terra-Coat L-21 12 oz plus PCNB-Terrazole granular 1 lb active per acre, and PCNB-Terrazole granular 1 lb active per acre. Isobac 20 was applied as an infurrow spray at planting at the rate of 6 oz per acre, as a spray over the cottonseed bed at ground cracking, and as a topical spray after emergence of the cotton seedlings. Delta Pine 16 aciddelinted cottonseed was planted June 25, 1969. Rhizoctonia inoculum was added in-furrow at planting time. Stand counts were taken of healthy appearing plants in 50 ft of row July 15. The results are as follows:

Treatment		ints ft row
Panagen 15 2 cz	129	a*
Isobac 20 in-furrow spray at planting 6 oz	179	a
Ischae 20 6 oz at ground cracking	192	ab
Isobac 20 6 cz post emergence	246	b
Panagen plus Vitavax 75W 8 oz PCNB-Terrazole granular	376	c .
1 lb active per acre	412	cd
Ceresan L 2 oz plus Demosan 65W 10 oz Terra-Chat L-21 12 cz, plus PCNB-Terrazole aranular	431	d
1 lb active per acre	476	d

^{*} Significant at 1% level.

These results are similar to previous trials in that Terra-Coat seed treatment plus PCNB-Terrazole granular, PCNB-Terrazole granular alone, or Ceresan L plus Demosan combination seed treatment were significantly better than the other treatments. Plants from seeds treated with Panogen plus Vitavax were next in effectiveness. Isobac performed poorly in this test and in two out of three cases was not significantly better than the check or no treatment.

Imperial Valley, 1969

Four growers trials were conducted during the spring of 1969 to compare single fungicide seed treatments. The seed treatments were as follows: Demosan 65W 10 oz, PCNB LST 12 oz, Panogen 15 2 oz, and no treatment. Rates of fungicides were per 100 lbs of untreated, acid-delinted Delta Pine 16 cottonseed. Disease incidence was low during these experiments and in only two trials were the responses significantly better than those from plots with no treatment. The

Brock trial was planted March 11 on a Holtville salty loam soil. The Borchard trial was planted March 27 on a Rositas very fine sandy loam soil. Plots were 200 ft long and replicated six times. The number of healthy plants per 100 ft of row was counted at the Brock ranch April 9 and the Borchard ranch April 22:

Treatment	Healthy plants per 100 ft row		
	Brock	Borchard	
Panogen 15 2 cz	914 a*	604 a	
PCNB LST 12 oz	7 95 a	554 a	
Demosan 65W 10 oz	646 b	591 a	
Check or no treatment	617 b	409 b	

^{*} Significant at 1% level

Panogen and PCNB were significantly better than Demosan or the check treatment at the Brock ranch. Panogen, PCNB, or Demosan seed treatment were significantly better than the check or no treatment at the Borchard ranch trial.

Meloland, 1969

Delta Pine 16 cotton seed was the variety used again in the spring trial at the Meloland Field Station. All fungicide rates are per 100 lbs of acid-delinted cotton seed or per-acre for the granular treatments. Treatments were as follows: No treatment, PCNB LST 12 oz, Terra-Coat L-21 12 oz, Vitavax 75W 8 oz, Ceresan L 2 oz plus Demosan 65W 10 oz, Demosan 65W 10 oz, and PCNB in-furrow granular 1 lb active. The plot was planted April 22 and treatments were 85 ft long and replicated six times. The number of healthy plants in 60 ft of row May 25: Healthy

Treatment		plants per 60 ft row	
Check cr no treatment	76	a*	
PCNB LST 12 oz	186	b	
PCNB in-furrow granular, 1 lb active	237	bc	
Terra-Coat L-21 12 oz	248	bc	
Vitavax 75W 8 oz	264	cd	
Ceresan L 2 oz plus Demosan 65W 10 oz	275	cd	
Demosan 65W 10 oz	328	d	

^{*} Significant at 1% level.

Vitavax, Demosan or a combination seed treatment of Ceresan plus Demosan were significantly better than all other treatments. Terra-Coat or PCNB in-furrow granular was intermediate in the number of healthy plants.

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Effects of SEMEN on

of

No differences in fertility and hatchability of turkey eggs were observed when hens were inseminated with semen extended with three different diluents tested, as compared with undiluted semen.

DILUTION OF TURKEY SEMEN has obvious economic advantages to the turkey breeder (as illustrated by the tom-hen ratio diagram). With a dilution program the semen from one male will inseminate three times the number of hens as in a program using undiluted semen. The more frequently the males are used within a week, the greater are the economic benefits from a dilution program with turkey breeder males.

Researchers in the Nebraska Experiment Station showed that turkey semen could be extended successfully in several experiments in 1954. Recently, Brown of the Ohio Experiment Station, and Lake of the Edinburgh Poultry Research Centre (using diluents widely different in composition), showed that dilution of turkey semen had no adverse effect upon fertility and hatchability of turkey eggs. At present, a diluent prepared by the Minnesota Turkey Grower's Association and sealed in 2-cc glass vials has been widely used in the field for the extension of turkey semen.

In this study, the objective was to determine whether these diluents (used at a ratio of one part semen to two parts diluent) would differ among themselves in terms of fertility and hatchability, after insemination into hens within one hour of collection and dilution. The formulae of the three diluents are shown in table 1.

three EXTENDERS reproduction turkeys

F. X. OGASAWARA · R. A. ERNST

Lake's and Brown's diluents were prepared aseptically and sealed in 2-cc glass vials (1.33 cc of diluent per vial). The Minnesota diluent was provided by Dr. E. Graham of the University of Minnesota in similar vials for use in the experiment. All diluents were stored in a refrigerator at $40^{\circ}F$ until the day of use.

One hundred twenty Bronze turkey females were randomly distributed into five treatment groups in four replicate blocks, so that full sisters appeared in all five treatments whenever possible. The hens were held in individual cages at the University of California experimental farm at Davis. The five treatments compared in this experiment are shown in table 2. The last two treatments are controls: one duplicating the total volume inseminated, and the other the actual number of spermatozoa inseminated.

Thirty-two Bronze males were held in flat bottom cages. The large number was for the purpose of insuring a surplus volume of semen, both for insemination and for semen quality testing. The collection, dilution, and insemination of semen was carried out within a one-hour period. The hens were inseminated on a biweekly schedule for a 13-week experimental period. The order of insemination was determined randomly on each date to eliminate any effect of time between collection of semen and insemination of the diluted semen.

After each insemination of the hens, the semen (both diluted and control samples) was evaluated for concentration and WITHOUT SEMEN DILUTION: 1 TOM PER 10 HENS.

WITH SEMEN DILUTION: 1 TOM PER 30 HENS.

WITH SEMEN DILUTION: 1 TOM PER 30 HENS.

for the percentage of live sperm. As shown in table 3, the concentration of spermatozoa in the diluted samples was proportionately lower than that of the whole semen.

Table 4 shows that egg production of the five treatment groups was reasonably uniform during this trial. Fertility was high, averaging 90 per cent in all groups for the duration of the trial. Table 4 also shows the per cent hatch of fertile eggs and of total eggs set, and again illustrates how closely these three treatments were grouped during the experimental period. Statistical analysis indicated that none of the means of the five treatment groups were significantly different from each other for egg production, fertile eggs, hatch of fertile eggs or hatch to total eggs set.

No differences in fertility and hatchability of eggs were observed when hens were inseminated with semen extended with the three diluents—or with undiluted semen—irrespective of dose used. It was concluded that any diluent (if osmotically balanced for turkey sperm cells) can be used as an extender if the collection, dilution and insemination procedures are confined to a one-hour interval.

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TABLE 1
DILUENT COMPOSITION FOR TURKEY SEMEN

Constituent	Lake's	Brown's	Minnesota
		gms per liter	
Mg Chloride	0.676		
K Citrate	1.280		
Na Acetate	8.511		
Na Glutamate	17.3 <i>5</i> 0		
Fructose	10.000		
Dextrose			2.000
Na Chloride		10.000	3.350
Di-Na Phosphate			1.185
K Di-Acid Phosphate			0.260
Ma Sulfate			0.500
K Chloride			2.200
Ca Chloride			0.500
Tris			1.670
Tricine			22.000
Peptone			20.000
Threonine			0.011
pH:	7.0	6.8	7.2

TABLE 2. EXPERIMENTAL DESIGN

Semen diluent	Dilution	Volume inseminated	
	semen:diluent	ml	
Lake's	1:2	0.03	
Minnesota	1:2	0.03	
Brown's	1:2	0.03	
None	None	0.03	
None	None	0.01	

TABLE 3. SEMEN QUALITY TESTS

S	cncentration	Live	e-dead values	
Sample	of sperm	Normal Abnormal		Dead
	billion/ml	%	%	%
Lake's diluent	2.5	72	20	8
Minnesota dilue	nt 2.3	75	17	8
Brown's diluent	2.5	75	16	9
Whole semen	9.6	64	23	13

TABLE 4. EGG PRODUCTION, FERTILITY, AND HATCHABILITY RESULTS

Semen: whole or diluted	Egg pro- duction	Fertile eggs	Hatch of fertile eggs	Hatch of total egg set
	%	%	%	%
Lake's	41	90	70	63
Minnesota	41	89	69	61
Brown's	38	89	65	59
Whole semen	39	93	71	65
⅓ semen	33	89	65	58
Average	38	90	68	61