

SEED GERMINATION IN CERTAIN NEW MEXICO RANGE GRASSES

CONTRIBUTIONS FROM THE HULL BOTANICAL LABORATORY 385

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(WITH FIVE FIGURES)

Introduction

Much work has been carried on in recent years in regard to the viability and germination of seeds of various plants. Seed-testing studies are highly important, because the results obtained influence or affect the work of the farmer, the floriculturist, the amateur gardener, and the rancher or ranger. These people cannot afford to plant seeds, expecting a 95 per cent germination and then perhaps having only a 50 per cent germination, due to poor selection of seed, to its immaturity, or to its adulteration with weed seeds.

Among the plants studied for germination the grasses have a prominent part, and this is especially true of the range grasses of the west. It is necessary for the rancher or ranger to know about what percentage of the seeds of the grasses covering his grazing lands he can expect to germinate, since he can then estimate the amount of grazing his lands will tolerate without becoming depleted. He must know whether his grasses propagate themselves vegetatively or by seed, or by both means.

Various studies have been made with reference to seed germination, and although they have no direct bearing upon the problem which the writer undertook, yet they gave ideas for some tests which were performed after the major problem was completed. The following are short summaries of the work of some of the investigators in the field of seed germination.

In general it is believed that the delay in after-ripening is due to the characters of the embryo and of the seed coat. Miss ECKERSON (6) found that there is a series of metabolic changes going on in the embryo during the period of after-ripening. At first the acidity is increased, and correlated with it is increased activity of catalase

and peroxidase. By treating the embryos with dilute acids such as hydrochloric, butyric, and acetic, the after-ripening period can be reduced very much. Those embryos which are treated increase their water-holding power, acidity, and amount of peroxidase more rapidly; and the oxidase appears sooner than in the untreated embryo.

PAMMEL and KING (14) planted mature and immature seeds in the fall and in the spring. In general, stratification in sand and freezing were favorable to germination. *Asclepias syriaca* showed 12 per cent germination, *Ambrosia psilostachya* 18 per cent, *Chenopodium album* 88 per cent, and *Xanthium canadense* 25 per cent.

PACK (13) discovered that the germination of non-after-ripened *Juniperus* seeds under ordinary conditions is very small, amounting to 1 per cent. These seeds are protected by a semi-permeable and thick coat which makes up 75 per cent by weight of the entire seed. Acids enter very slowly, he found, while bases, silver, and mercury salts enter rapidly. PACK thinks that while the coat may act as a protection against fungal attack, and may prevent water-imbibed seeds from expanding and bursting the tissues before after-ripening is accomplished, it takes little or no part in the dormancy of after-ripening of the seed. He was unable to force the germination of non-after-ripened *Juniperus* seeds by high temperature, alternate temperatures, wounding, warm bath, dry air, removal of coats; or by treatment with hydrogen peroxide, mercuric chloride, ether, carbon dioxide, oxygen, light, soil, dilute acids, dilute bases, nitrates, sulphates, or strong acids. Freezing and thawing as such have no forcing action on germination of the *Juniperus* seeds, neither do they hasten after-ripening. They bring about chemical changes in the seed, but these changes are different from those occurring during after-ripening. When seeds are about ready to germinate, PACK found that they are very sensitive and are killed by exposure to 5° C. The *Juniperus* seed has a dormant embryo that must be after-ripened before germination.

Mrs. DAVIS (4) found the sterilization of the naked seeds of *Cornus florida* difficult, due to the fact that the inner testa is very thin or the endosperm rich in food and the seeds are frequently infested with molds. She concludes that the pericarp, the outer testa, and the inner testa take no part in causing the delay; the moisture

intake with the pericarp and testas intact is as high as in seeds with these removed. After-ripened seeds break the pericarp readily. Delay is not caused by an immature embryo, as it is well differentiated several weeks prior to shedding. After-ripening of the dormant seed is favored by low temperatures, 0–5° C.

Rubus seeds vary according to species in the time required to weaken the coat with sulphuric acid. After the carbonized coats have been removed from the seeds they must be carefully sterilized, since treated seeds are very susceptible to molds. The delay in *Sphaeralcea remota* is due to the impermeable cuticle forming the outer layer of the seed coat. After this coat has been subjected to a 2–3 hour treatment with sulphuric acid it becomes permeable. Chipping slightly helps, as this method makes it possible for water to enter the seed. Mrs. DAVIS found that untreated seeds failed to swell or germinate, but that may have been due to the fact that the seeds were gathered late in the season. Treated seeds germinated and grew for a time in distilled water, and all seedlings showed remarkable vigor.

CROCKER (3) claims that in *Xanthium canadense* delayed germination is generally due to the seed coat rather than to the embryo. In the upper cockle-bur seed the delay is due to the exclusion of oxygen by the seed coat. No germination appeared in *Iris* seeds because the cap and endosperm stopped the absorption of water before the needed amount was obtained by the embryo. Those seed coats which exclude water are better for causing delay than those which exclude oxygen, because there is less respiration. The length of delay is due in nature to the persistence of the seed coats. In the case of *Xanthium*, the bur helps in causing the upper seed to germinate later. Seed coats reduce the oxygen supply, especially in the upper seed. High temperature brings about the germination of the upper seeds with the coats intact by raising the respiration ratio, which increases the rate of diffusion of oxygen through the seed coat.

DUVEL (5) has discovered that the factors affecting the vitality of the seed are maturity, weather conditions at time of harvesting, methods of harvesting, and curing. Immature seeds sown soon after gathering usually germinate readily, but if they are stored they soon lose their vitality. Seeds which are harvested in damp rainy weather are much weaker in vitality. By special care the life of the seed once

injured may be prolonged. Since moisture affects the longevity of seeds, they must be kept in a dry place where the temperature is low. DUVEL found that seeds treated in a sulphuric bath or in a vacuum usually showed delayed germination because the seed coat has hardened. In order to keep the vitality of the seed, it is better to have as little respiration as possible. Respiration brings about a chemical activity in the cells, which causes energy to be transformed, resulting eventually in the death of the seed. Respiration in the light is the same as in the dark if moisture and temperature conditions are the same.

FAWCETT (8) worked with 92 samples of weed seeds, representing 52 species. The seeds were collected in September, October, and November. Fifty seeds of each sample were placed in sand in boxes under the benches in the greenhouse. Every month from November to May this was repeated. Samples were also placed in sacks inside of a wooden box and a thin layer of sand placed around them. The boxes were then sunk in the ground so that just the top was exposed. Comparisons were made with the samples kept indoors. In April both lots were planted outdoors. FAWCETT concludes that weed seeds with thick seed coats require a more or less extended period of rest after maturity. Mustard and pepper grass seeds require little time for rest. Drying out weakens the vitality of nearly all weed seeds, and exposure to the natural periods for best seed germination, fall and spring, increases the power of germination.

HIMMEL (11) also believes that low percentage of germination for honey locust is due to the seed coats, for when the seeds had been treated with concentrated sulphuric acid the germination percentage increased very much. The older the wheat seeds the lower the germination percentage, but *Amaranthus* seeds displayed greater germination in the older seeds. The life of a dandelion seed is less than 8 years, but even the old seeds show good catalase activity. *Typha* seeds germinate better if they are pricked. Only very dilute acids and bases have any forcing effect on *Typha* seeds.

MISS EVANS (7) found that in after-ripened seeds with coats untreated, the restricting effect of the coats showed particularly at low temperatures 8–10° and 11.6° C., and again at high temperatures, 42° for Washington seeds and 46.1° for Indiana seeds. In both

cases these effects can be lessened by treating the coats with H_2SO_4 , or abrading them with sand.

ATWOOD (1) found that there was less delay in germination in *Avena fatua* after the shell coats had been removed. Restriction of the oxygen supply by the seed coat acts as a limiting factor in germination. These seeds do not seem to be affected by light during germination. ATWOOD concludes that after-ripening occurs with the drying of the seed, but independent of the water content, as air-dried seeds soon after harvest yield lower germinative percentages than seeds of similar moisture content the following spring. Exclusion of water by the true seed coat does not explain after-ripening according to ATWOOD.

Material and methods

The primary purpose of this investigation was to discover the percentage of germination of the 1926 seeds of the following grasses:

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|-----------------------------------|-----------------------------------|
| 1. <i>Hilaria mutica</i> | 10. <i>Bouteloua gracilis</i> |
| 2. <i>H. mutica</i> | 11. <i>Bouteloua eriopoda</i> |
| 3. <i>Muhlenbergia porteri</i> | 12. <i>Sporobolus giganteus</i> |
| 4. <i>Sporobolus airoides</i> | 13. <i>Aristida longiseta</i> |
| 5. <i>Aristida purpurea</i> | 14. <i>Sporobolus flexuosus</i> |
| 6. <i>Scleropogon brevifolius</i> | 15. <i>Aristida longiseta</i> |
| 7. <i>Bouteloua eriopoda</i> | 16. <i>Bouteloua curtipendula</i> |
| 8. <i>Muhlenbergia arenicola</i> | 17. <i>Sporobolus auriculatus</i> |
| 9. <i>Sporobolus cryptandrus</i> | |

Studies of these grasses have been made in previous years at the government laboratories in Washington, D.C., but the 1926 seeds were sent to the Hull Laboratory of the University of Chicago for experimentation by the writer. The seeds were collected by R. S. CAMPBELL on the Jornada Range Reserve Station at Las Cruces, New Mexico, and through him the following seeds from 1925 were obtained:

- | | |
|--------------------------------|-----------------------------------|
| 20. <i>Sporobolus airoides</i> | 24. <i>Epicampes emerslyei</i> |
| 21. <i>S. auriculatus</i> | 25. <i>Muhlenbergia arenicola</i> |
| 22. <i>S. cryptandrus</i> | 26. <i>Panicum obtusum</i> |
| 23. <i>S. flexuosus</i> | |

The seeds were still in the glumes on the spikes, hence it was necessary to separate not only the florets from the spikelets, but also

to remove the palea and lemma from the seeds. The seeds for the tests were selected at random, so that the results would be as representative as possible. Since some of the samples contained relatively few seeds, it was found necessary to use smaller amounts of seeds for some tests. In all cases the seeds were soaked for one half-hour in a 0.25 per cent solution of uspulun and then rinsed (washed) in distilled water twice for 15 minute periods. The blotting paper was also treated with the uspulun solution, and the Petri dishes containing cotton and filter paper were sterilized by autoclaving. These precautions were taken to insure the seeds against black mold attack. Only in a few instances were the seeds attacked by molds.

All seeds were tested at 25° C. in the moist chamber germinating oven in blotters and in Petri dishes. Those which did not do so well at 25° were tried at 35° C. Each species was tested for 100 per cent germination. In the preliminary tests, 10 seeds of each species were used. In some instances the lemma and the palea were not removed in the first tests, hence it was not certain that there were seeds present. Afterward it was made a point that all protective bracts be removed from the seeds. Exceptions were made in the case of the *Bouteloua* grasses, in which the writer could find no seeds; hence the entire florets were used in hopes that some might contain seeds.

The *Aristida* seeds were tested for germination in the light, as they were supposed to show better germination results there than in the dark. The *Sporobolus* seeds were treated in several ways. The seed coats of nos. 9, 12, 14 were pricked, and the seeds shaken for 4, 6, and 9 hours in bottles containing coarse white sand, in order to injure the seed coats, thus hastening germination through the more easy entrance of water into the seed. Some of the seeds of nos. 9, 12, 14, which had been shaken for various lengths of time, were planted in sandy loam and kept at room temperature to see whether they would germinate more readily after having their seed coats bruised. Nos. 9, 12, 14, 22, 23 were soaked in distilled water for a period of 4 days and then for one of 9 days. These seeds were then placed in the 5° C. oven for 7 days and 21 days, and then in the 25° C. germinating oven. Nos. 4 and 20 were treated with varying solutions of CaCO₃. Ten seeds of all the grasses, excepting the *Bouteloua* species, were planted in sandy loam.

TABLE I A
TABULAR DESCRIPTION OF SEEDS COLLECTED BY R. S. CAMPBELL

SEED NO.	SCIENTIFIC NAME	COMMON NAME	GEOGRAPHIC RANGE	PLACE OF COLLECTION	DATE OF COLLECTION (1926)
1.	<i>Hilaria mutica</i>	Tabosa grass	Western Texas to southern Arizona and adjacent Mexico	1 1/4 m. southwest of Red Lake Well	Sept. 19
2.	<i>Hilaria mutica</i>	Tabosa grass	Western Texas to southern Arizona and adjacent Mexico	1 m. south of Middle Well	Sept. 1
3.	<i>Muhlenbergia porteri</i>	Porter's bush grama	Colorado and western Texas to California and Mexico	2 m. south of Taylor Well
4.	<i>Sporobolus atroideus</i>	Alkali sacaton	Washington and Nebraska to California and New Mexico	1 1/4 m. northeast of Hdqrs.	Aug. 31
5.	<i>Aristida purpurea</i>	Purple three-awn	Southwestern U.S. to southern Mexico	1 1/2 m. northeast of Hdqrs.	Aug. 31
6.	<i>Scleropogon brevifolius</i>	Burro grass	Arizona and western Texas to Mexico and South America	3/4 m. northeast of Hdqrs.	Aug. 31
7.	<i>Bouteloua eriopoda</i>	Black grama grass	Arizona and western Texas to Mexico	1 1/4 m. northeast of Hdqrs.	Aug. 31
8.	<i>Muhlenbergia arenicola</i>	Ring muhlenbergia	Colorado and Kansas to Texas and New Mexico	1 1/2 m. northeast of Hdqrs.	Aug. 31
9.	<i>Sporobolus cryptandrus</i>	Sand grass	Washington and Maine to Arizona and Texas	1 1/4 m. northeast of Hdqrs.	Aug. 31
10.	<i>Bouteloua gracilis</i>	Blue grama grass	Manitoba to Mexico, and even to South America	St. Nicholas Canyon enclosure	Sept. 1
11.	<i>Bouteloua eriopoda</i>	Black grama grass	Arizona and western Texas to Mexico	Enclosure no. 10	Nov. 9
12.	<i>Sporobolus giganteus</i>	Gigantic sand grass	Southern New Mexico	1 1/4 m. northeast of Hdqrs.	Sept. 19
13.	<i>Aristida longiseta</i>	Red three-awn grass	Southwestern U.S. and New Mexico	Aristida enclosure	Sept. 16
14.	<i>Sporobolus flexuosus</i>	Wide-panicked grass	Nevada to Texas and Mexico	1 1/4 m. northeast of Hdqrs.	Aug. 31
15.	<i>Aristida longiseta</i>	Red three-awn grass	Southwestern U.S. and northern Mexico	1 1/4 m. northeast of Hdqrs.	Aug. 31
16.	<i>Bouteloua curtipendula</i>	Side oats grama grass	Canada to New Jersey, California, and Mexico	Lion Den Canyon	Sept. 1
17.	<i>Sporobolus auriculatus</i>	Dwarf dropseed	Western Texas to southern New Mexico	3/4 m. north of Hdqrs.	Aug. 31

TABLE IB
TABULAR DESCRIPTION OF SEEDS

SEED NO.	SCIENTIFIC NAME	ALTITUDE OF PLACE OF COLLECTION (FEET)	TYPE OF SOIL	RAINFALL				PERCENT-AGE DISSEMINATION	PERCENT-AGE GERMINATION
				SEASONAL		ANNUAL			
				Inches	Zone	Inches	Zone		
1.	<i>Hilaria mutica</i>	4200	Heavy clay	4.82	1	14.67	1	34	
2.	<i>Hilaria mutica</i>	4200	Low swag heavy clay	7.51	4	18.29	5	91	
3.	<i>Muhlenbergia porteri</i>	4050	Gravelly clay-sand	10.24	6	19.48	5	75	
4.	<i>Sporobolus airoides</i>	4200	Clay loam	9.64	6	17.42	4	92	
5.	<i>Aristida purpurea</i>	4200	Sandy loam	9.64	6	17.42	4	60	
6.	<i>Scleropogon brevifolius</i>	4200	Clay loam	8.53	5	17.42	4	46	
7.	<i>Bouteloua eriopoda</i>	4200	Sandy loam	9.64	6	17.42	4	60	
8.	<i>Muhlenbergia arenicola</i>	4200	Clay loam	9.64	6	17.42	4	60	
9.	<i>Sporobolus cryptandrus</i>	4200	Sandy loam	9.64	6	17.42	4	75	
10.	<i>Bouteloua gracilis</i>	5600	Gravelly loam	11.23	8	21.17	7	5	
11.	<i>Bouteloua eriopoda</i>	4050	Gravelly loam	7.65	4	17.42	4	97	
12.	<i>Sporobolus giganteus</i>	4200	Sandy loam	9.64	6	17.42	4	17	
13.	<i>Aristida longisetia</i>	4300	Very sandy loam	5.73	2	18.53	2	25	
14.	<i>Sporobolus flexuosus</i>	4200	Sandy loam	9.64	6	17.42	4	30	
15.	<i>Aristida longisetia</i>	4200	Sandy loam	9.64	6	17.42	4	53	
16.	<i>Bouteloua curtipendula</i>	4800	Gravelly loam	11.23	7	21.17	6	10	
17.	<i>Sporobolus auriculatus</i>	4100	Clay loam	8.53	5	17.42	4	100	

TABLE II

PRELIMINARY TEST ON BLOTTING PAPER, 25° C.; JANUARY 19-FEBRUARY 18, 1927
(31 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
1.....	<i>Hilaria mutica</i>	10	1/27	10
2.....	<i>Hilaria mutica</i>	10	1/27	60
3.....	<i>Muhlenbergia porteri</i>	10	1/27	10
4.....	<i>Sporobolus airoides</i>	10	1/24	40
5.....	<i>Aristida purpurea</i>	9	1/24	33.3
6.....	<i>Scleropogon brevifolius</i>	10	2/3	20
7.....	<i>Bouteloua eriopoda</i>	10*
8.....	<i>Muhlenbergia arenicola</i>	10	2/3	30
9.....	<i>Sporobolus cryptandrus</i>	10	2/10	30
10.....	<i>Bouteloua gracilis</i>	10*
11.....	<i>Bouteloua eriopoda</i>	10*
12.....	<i>Sporobolus giganteus</i>	10	2/10	50
13.....	<i>Aristida longiseta</i>	10	Moldy
14.....	<i>Sporobolus flexuosus</i>	10	2/15	80
15.....	<i>Aristida longiseta</i>	10	1/24	50
16.....	<i>Bouteloua curtipendula</i>	10	Moldy
17.....	<i>Sporobolus auriculatus</i>	10	1/27	20

* Empty lemmas.

TABLE III

PRELIMINARY TESTS ON BLOTTING PAPER, 25° C.; FEBRUARY 4-25, 1927 (22 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
1.....	<i>Hilaria mutica</i>	10	2/7	20
2.....	<i>Hilaria mutica</i>	10	2/7	100
3.....	<i>Muhlenbergia porteri</i>	10	2/7	80
4.....	<i>Sporobolus airoides</i>	10	2/15	20
5.....	<i>Aristida purpurea</i>	10	2/10	70
6.....	<i>Scleropogon brevifolius</i>	10	2/25	10 (9 moldy)
7.....	<i>Bouteloua eriopoda</i>	10*
8.....	<i>Muhlenbergia arenicola</i>	10	2/10	30
9.....	<i>Sporobolus cryptandrus</i>	10	2/15	60
10.....	<i>Bouteloua gracilis</i>	10*
11.....	<i>Bouteloua eriopoda</i>	10*
12.....	<i>Sporobolus giganteus</i>	10	2/25	40
13.....	<i>Aristida longiseta</i>	10	2/7	30
14.....	<i>Sporobolus flexuosus</i>	10	2/25	30
15.....	<i>Aristida longiseta</i>	10	2/7	40
16.....	<i>Bouteloua curtipendula</i>	10	2/7	100
17.....	<i>Sporobolus auriculatus</i>	10	2/7	20

* Empty lemmas.

TABLE IV

PRELIMINARY TEST IN PETRI DISH, 25° C.; JANUARY 27—FEBRUARY 25 (30 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
1.....	<i>Hilaria mutica</i>	10	2/1	40
2.....	<i>Hilaria mutica</i>	10	1/29	100
3.....	<i>Muhlenbergia porteri</i>	10	2/1	60
4.....	<i>Sporobolus airoides</i>	10	1/29	70
5.....	<i>Aristida purpurea</i>	10	1/29	80
6.....	<i>Scleropogon brevifolius</i>	10	Moldy
7.....	<i>Bouteloua eriopoda</i>	10*
8.....	<i>Muhlenbergia arenicola</i>	10	2/1	40
9.....	<i>Sporobolus cryptandrus</i>	10	1/22	40
10.....	<i>Bouteloua gracilis</i>	10*
11.....	<i>Bouteloua eriopoda</i>	10*
12.....	<i>Sporobolus giganteus</i>	10	1/22	40
13.....	<i>Aristida longiseta</i>	10	2/1	10
14.....	<i>Sporobolus flexuosus</i>	10	2/22	70
15.....	<i>Aristida longiseta</i>	10	2/1	80
16.....	<i>Bouteloua curtipendula</i>	10	2/3	100
17.....	<i>Sporobolus auriculatus</i>	9	2/1	100

* Empty lemmas.

TABLE V

PRELIMINARY TEST IN PETRI DISH, 35° C.; FEBRUARY 2—16 (14 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
1.....	<i>Hilaria mutica</i>	10	2/7	40
2.....	<i>Hilaria mutica</i>	Not tried
3.....	<i>Muhlenbergia porteri</i>	Not tried
4.....	<i>Sporobolus airoides</i>	Not tried
5.....	<i>Aristida purpurea</i>	Not tried
6.....	<i>Scleropogon brevifolius</i>	10	0
7.....	<i>Bouteloua eriopoda</i>	10*	0
8.....	<i>Muhlenbergia arenicola</i>	5	0
9.....	<i>Sporobolus cryptandrus</i>	10	0
10.....	<i>Bouteloua gracilis</i>	10*	0
11.....	<i>Bouteloua eriopoda</i>	10*	0
12.....	<i>Sporobolus giganteus</i>	10	0
13.....	<i>Aristida longiseta</i>	10	2/7	10
14.....	<i>Sporobolus flexuosus</i>	10	0
15.....	<i>Aristida longiseta</i>	Not tried
16.....	<i>Bouteloua curtipendula</i>	10	2/7	100
17.....	<i>Sporobolus auriculatus</i>	Not tried

* Empty lemmas.

TABLE VI

PRELIMINARY TEST IN PETRI DISH, 25° C.; FEBRUARY 12-MARCH 15 (32 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
1.....	<i>Hilaria mutica</i>	10	2/15	20
2.....	<i>Hilaria mutica</i>	10	2/15	80
3.....	<i>Muhlenbergia porteri</i>	10	2/15	10
4.....	<i>Sporobolus airoides</i>	10	2/17	90
5.....	<i>Aristida purpurea</i>	10	2/15	40
6.....	<i>Scleropogon brevifolius</i>	10	2/15	40
7.....	<i>Bouteloua eriopoda</i>	10*
8.....	<i>Muhlenbergia arenicola</i>	4	2/15	50
9.....	<i>Sporobolus cryptandrus</i>	10	2/15	100
10.....	<i>Bouteloua gracilis</i>	10*
11.....	<i>Bouteloua eriopoda</i>	10*
12.....	<i>Sporobolus giganteus</i>	10	2/15	100
13.....	<i>Aristida longiseta</i>	10	2/15	10
14.....	<i>Sporobolus flexuosus</i>	10	2/15	60
15.....	<i>Aristida longiseta</i>	10	2/15	60
16.....	<i>Bouteloua curtipendula</i>	10	2/15	100
17.....	<i>Sporobolus auriculatus</i>	9	2/15	33.3

* Empty lemmas.

TABLE VII

100 PER CENT GERMINATION TEST IN PETRI DISH, 25° C.; FEBRUARY 21-MARCH 16 (24 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
1.....	<i>Hilaria mutica</i>	50	2/28	34
2.....	<i>Hilaria mutica</i>	100	2/25	91
3.....	<i>Muhlenbergia porteri</i>	20	2/28	75
4.....	<i>Sporobolus airoides</i>	100	2/25	92
5.....	<i>Aristida purpurea</i>	100	2/25	60
6.....	<i>Scleropogon brevifolius</i>	50	2/28	48
7.....	<i>Bouteloua eriopoda</i>	100*
8.....	<i>Muhlenbergia arenicola</i>	20	2/25	60
9.....	<i>Sporobolus cryptandrus</i>	100	2/25	75
10.....	<i>Bouteloua gracilis</i>	100*
11.....	<i>Bouteloua eriopoda</i>	100*
12.....	<i>Sporobolus giganteus</i>	100	2/25	97
13.....	<i>Aristida longiseta</i>	100	2/25	17
14.....	<i>Sporobolus flexuosus</i>	100	2/25	42
15.....	<i>Aristida longiseta</i>	100	2/25	53
16.....	<i>Bouteloua curtipendula</i>	75	2/28	98.6
17.....	<i>Sporobolus auriculatus</i>	25	2/28	100

* Empty lemmas.

TABLE VIII

PRELIMINARY TEST ON BLOTTING PAPER, 25° C.; JANUARY 19—FEBRUARY 18 (31 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
20.....	<i>Sporobolus airoides</i>	10	1/29	100
21.....	<i>Sporobolus auriculatus</i>	10*
22.....	<i>Sporobolus cryptandrus</i>	10	2/10	80
23.....	<i>Sporobolus flexuosus</i>	10	2/10	60
24.....	<i>Epicampes emersleyi</i>	10*
25.....	<i>Muhlenbergia arenicola</i>	10	2/1	30

* Empty lemmas.

TABLE IX

PRELIMINARY TEST IN PETRI DISH, 25° C.; JANUARY 27—FEBRUARY 25 (30 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
20.....	<i>Sporobolus airoides</i>	10	2/1	100
21.....	<i>Sporobolus auriculatus</i>	Empty lemmas
22.....	<i>Sporobolus cryptandrus</i>	10	2/22	70
23.....	<i>Sporobolus flexuosus</i>	10	2/22	80
24.....	<i>Epicampes emersleyi</i>	Empty lemmas
25.....	<i>Muhlenbergia arenicola</i>	10	2/1	40
26.....	<i>Panicum obtusum</i>	10	2/7	30

TABLE X

PRELIMINARY TEST IN PETRI DISH, 35° C.; FEBRUARY 2—16 (14 DAYS)

No.	NAME	No. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
20.....	<i>Sporobolus airoides</i>	Not tried
21.....	<i>Sporobolus auriculatus</i>	5	2/10	20
22.....	<i>Sporobolus cryptandrus</i>	10
23.....	<i>Sporobolus flexuosus</i>	10
24.....	<i>Epicampes emersleyi</i>	Empty lemmas
25.....	<i>Muhlenbergia arenicola</i>	9	2/7	77.8
26.....	<i>Panicum obtusum</i>	10

TABLE XI

100 PER CENT GERMINATION TEST IN PETRI DISH, 25° C.; FEBRUARY 21-MARCH 16
(24 DAYS)

No.	NAME	NO. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
20.....	<i>Sporobolus airoides</i>	100	2/25	92
21.....	<i>Sporobolus auriculatus</i>	6	2/25	100
22.....	<i>Sporobolus cryptandrus</i>	20	2/25	95
23.....	<i>Sporobolus flexuosus</i>	40	2/25	92.5
24.....	<i>Epicamps emersleyi</i>	Not tried
25.....	<i>Muhlenbergia arenicola</i>	40	2/25	90
26.....	<i>Panicum obtusum</i>	100	2/28	21

TABLE XII

GERMINATION IN LIGHT IN PETRI DISH, ROOM TEMPERATURE, ROOM LIGHT;
MARCH 1-16 (16 DAYS)

No.	NAME	NO. USED	DATE OF FIRST GERMINATION	FINAL GERMINATION PERCENTAGE
5.....	<i>Aristida purpurea</i>	10	3/5	40
13.....	<i>Aristida longiseta</i>	10	3/5	20
15.....	<i>Aristida longiseta</i>	10	3/5	80

TABLE XIII

SEEDS SOAKED IN DISTILLED H₂O 4 DAYS, THEN IN GERMINATING OVEN AT 25° C.
FOR 2 WEEKS

No.	NAME	NO. USED	GERMINATION DURING SOAKING	GERMINATION AFTER SOAKING	PERCENTAGE GERMINATION
9.....	<i>Sporobolus cryptandrus</i>	15	0	0	0
12.....	<i>Sporobolus giganteus</i>	15	0	0	6.6
14.....	<i>Sporobolus flexuosus</i>	15	0	0	0
20.....	<i>Sporobolus airoides</i>	40	34	0	85
22.....	<i>Sporobolus giganteus</i>	15	0	0	0
23.....	<i>Sporobolus flexuosus</i>	15	0	0	0

TABLE XIV

SEEDS SOAKED IN DISTILLED H₂O 10 DAYS, THEN IN GERMINATING OVEN AT 25° C.
FOR 2 WEEKS

No.	NAME	NO. USED	GERMINATION DURING SOAKING	GERMINATION AFTER SOAKING	PERCENTAGE GERMINATION
9.....	<i>Sporobolus cryptandrus</i>	15	0	2	13.3
12.....	<i>Sporobolus giganteus</i>	15	2	0	13.3
14.....	<i>Sporobolus flexuosus</i>	15	1	0	6.6
20.....	<i>Sporobolus airoides</i>	15	12	0	80
22.....	<i>Sporobolus giganteus</i>	15	0	1	6.6
23.....	<i>Sporobolus flexuosus</i>	15	1	0	6.6

TABLE XV

SEEDS SHAKEN IN COARSE WHITE SAND, THEN IN 25° C. GERMINATING
OVEN FOR 2 WEEKS

No.	NAME	NO. USED	NO. OF HOURS SHAKEN	NO. OF DAYS SOAKED	PERCENTAGE GERMINATION
9.....	<i>Sporobolus cryptandrus</i>	10	6	5	10
12.....	<i>Sporobolus giganteus</i>	10	6	5	10
14.....	<i>Sporobolus flexuosus</i>	10	6	5	10

TABLE XVI

SEEDS SHAKEN IN COARSE WHITE SAND, THEN PLACED IN SANDY LOAM

No.	NAME	NO. USED	NO. OF HOURS SHAKEN	PERCENTAGE GERMINATION	NO. OF HOURS SHAKEN	PERCENTAGE GERMINATION
9.....	<i>Sporobolus cryptandrus</i>	10	6	0	9	0
12.....	<i>Sporobolus giganteus</i>	10	6	0	9	0
14.....	<i>Sporobolus flexuosus</i>	10	6	0	9	0

TABLE XVII
SEEDS IN GERMINATING OVEN AT 25° C. FOR 21 DAYS

No.	NAME (10 SEEDS OF EACH SPECIES USED)	PERCENTAGE GERMINATION
Without shaking or soaking (control)		
9.....	<i>Sporobolus cryptandrus</i>	10
12.....	<i>Sporobolus giganteus</i>	20
14.....	<i>Sporobolus flexuosus</i>	10
Shaken four hours		
9.....	<i>Sporobolus cryptandrus</i>	0
12.....	<i>Sporobolus giganteus</i>	0
14.....	<i>Sporobolus flexuosus</i>	0
Shaken six hours		
9.....	<i>Sporobolus cryptandrus</i>	10
12.....	<i>Sporobolus giganteus</i>	10
14.....	<i>Sporobolus flexuosus</i>	10
Shaken nine hours		
9.....	<i>Sporobolus cryptandrus</i>	10
12.....	<i>Sporobolus giganteus</i>	20
14.....	<i>Sporobolus flexuosus</i>	10

TABLE XVIII

No.	NAME	No. USED	PERCENTAGE GERMINATION	
			5° C. oven 7 days	25° C. oven 21 days
9.....	<i>Sporobolus cryptandrus</i>	15	0	0
12.....	<i>Sporobolus giganteus</i>	15	13.3	20
14.....	<i>Sporobolus flexuosus</i>	15	0	0
22.....	<i>Sporobolus cryptandrus</i>	15	0	0
23.....	<i>Sporobolus flexuosus</i>	15	0	0

TABLE XIX

SEEDS TREATED WITH CaCO_3 SOLUTIONS INSTEAD OF DISTILLED H_2O

No.	NAME	No. used	PERCENTAGE GERMINATION			
			0.5 per cent CaCO_3	1 per cent CaCO_3	5 per cent CaCO_3	10 per cent CaCO_3
4....	<i>Sporobolus airoides</i> (1926)	10	30	70	0	0
20....	<i>Sporobolus airoides</i> (1925)	15	87	80	0	0

TABLE XX

NO SEED SELECTION MADE; GLUMES PLACED IN PETRI DISHES
AND THEN IN 25° C. OVEN

No.	NAME	RESULTS
7.....	<i>Bouteloua eriopoda</i>	No signs of germination
10.....	<i>Bouteloua gracilis</i>	No signs of germination
11.....	<i>Bouteloua eriopoda</i>	No signs of germination

TABLE XXI

SEEDS PLANTED IN SANDY LOAM AND KEPT AT ROOM TEMPERATURE (70° F.)
FEBRUARY 28, 1927

No.	NAME	NO. USED	DATE OF FIRST PLANT
1.....	<i>Hilaria mutica</i>	10
2.....	<i>Hilaria mutica</i>	10	3/16
3.....	<i>Muhlenbergia porteri</i>	10
4.....	<i>Sporobolus airoides</i>	10
5.....	<i>Aristida purpurea</i>	10
6.....	<i>Scleropogon brevifolius</i>	6	5/4
8.....	<i>Muhlenbergia arenicola</i>	10
9.....	<i>Sporobolus cryptandrus</i>	10	5/5
12.....	<i>Sporobolus giganteus</i>	10	4/13
13.....	<i>Aristida longiseta</i>	10
14.....	<i>Sporobolus flexuosus</i>	10	4/14
15.....	<i>Aristida longiseta</i>	10
16.....	<i>Bouteloua curtipendula</i>	10
17.....	<i>Sporobolus auriculatus</i>	10

TABLE XXII
PRECIPITATION JORNADA RANGE RESERVE, 1926

STATION	JANU- ARY	FEBRU- ARY	MARCH	APRIL	MAY	JUNE	JULY	AUGUST	SEPTEM- BER	OCTO- BER	NOVEM- BER	DECEM- BER	SEASON- AL	ANNUAL
Headquarters.....	0.49	0.05	1.40	0.48	2.43	0.06	4.95	0.38	3.20	2.32	0.04	1.62	8.53	17.42
Midwell.....	0.56	0.01	2.28	0.59	3.09	0.01	3.21	1.06	3.24	2.23	0.04	1.97	7.51	18.29
Red Lake.....	0.49	0.01	1.91	0.74	2.97	0.02	1.85	0.42	2.55	1.75	0.02	1.94	4.82	14.67
Road Tank.....	0.01	0.01	1.82	0.67	1.71	0.00	5.19	0.22	3.91	2.37	0.06	1.78	9.32	18.35
Ropes.....	0.84	0.12	2.65	0.88	2.71	0.40	2.52	1.09	4.11	2.36	0.00	2.00	7.72	19.68
St. Nicholas.....	0.90	0.10	2.02	0.53	1.64	0.03	6.07	1.20	3.95	2.67	0.02	2.03	11.23	21.17
South Well.....	0.53	0.00	1.57	0.91	2.09	T	4.01	0.25	3.48	2.61	0.01	1.84	7.74	17.30
Stuart Well.....	0.55	0.00	1.97	0.71	1.93	T	4.75	0.25	2.78	2.66	0.04	1.73	7.78	17.37
Ragged Well.....	0.43	0.00	1.96	1.00	2.21	0.00	6.30	1.16	2.78	2.24	0.00	1.31	10.24	19.48
West Well.....	0.55	0.00	1.66	0.71	1.98	T	2.48	1.66	4.81	2.53	0.12	2.03	8.95	19.53
Period Study.....	1.62	0.51	2.00	0.01	5.42	0.52	3.70	2.52	0.00	1.89	9.64
Enclosure no. 1.....	1.51	0.68	2.45	0.00	4.00	0.25	3.64	2.35	0.00	2.05	7.89
Enclosure no. 10.....	1.37	0.88	1.94	0.02	3.81	0.31	3.53	2.34	0.00	2.00	7.65
Enclosure no. 2.....	2.11	0.47	4.16	2.43	0.01	1.78	6.74
Aristida.....	2.77	0.42	2.54	0.89	0.79	2.01	5.73
Brown Tank.....	5.76	0.04	3.21	2.30	0.03	1.36	9.01
Sand Hills.....	5.24	0.39	3.15	2.09	0.00	1.98	8.68
New Well.....	5.76	0.81	3.58	1.84	0.40	1.95	10.15
Average.....	0.59	0.03	1.92	0.72	2.28	0.06	4.23	0.60	3.46	2.25	0.13	1.85	8.29	18.23

TABLE XXIII

COMPARISON OF SEASONAL RAINFALL FOR (A) *HILARIA MUTICA* NO. 1 AND NO. 2;
(B) *ARISTIDA LONGISETA* NO. 13 AND NO. 15*

PRECIPITATION (INCHES)			
(a) <i>Hilaria mutica</i> , no. 1		<i>Hilaria mutica</i> , no. 2	
July.....	1.85	July.....	3.21
August.....	0.42	August.....	1.06
September (19 days).....	1.01		
Total.....	3.88	Total.....	4.27
(b) <i>Aristida longiseta</i> , no. 13		<i>Aristida longiseta</i> , no. 15	
July.....	2.77	July.....	5.42
August.....	0.42	August.....	0.52
September (16 days).....	1.52		
Total.....	4.71	Total.....	5.94

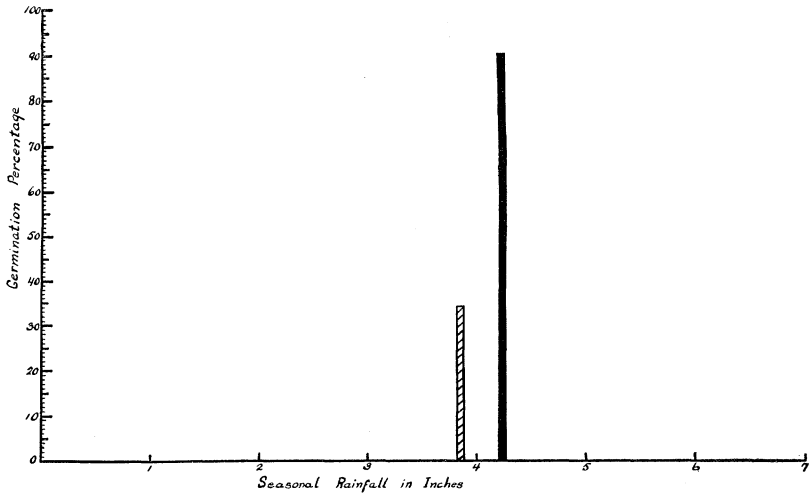
* See figs. 1, 2, 3.

TABLE XXIV

SEASONAL PRECIPITATION ON JORNADA RANGE RESERVE JULY, AUGUST,
AND SEPTEMBER, 1926

No.	STATION	PRECIPITATION (INCHES)	RANGE (INCHES)
1.....	Red Lake	4.82	Under 5
2.....	Aristida Enclosure	5.73	5-6
3.....	Enclosure no. 2	6.74	6-7
4.....	Middle Well	7.51	7-8
5.....	Enclosure no. 10	7.65	7-8
6.....	Ropes	7.72	7-8
7.....	South Well	7.74	7-8
8.....	Stuart Well	7.78	7-8
9.....	Enclosure no. 1	7.89	7-8
10.....	Headquarters	8.53	8-9
11.....	Sand Hills	8.68	8-9
12.....	West Well	8.95	8-9
13.....	Brown Tank	9.01	9-10
14.....	Road Tank	9.32	9-10
15.....	Period Study	9.64	9-10
16.....	New Well	10.15	10-11
17.....	Ragged Tank	10.24	10-11
18.....	St. Nicholas	11.23	Over 11

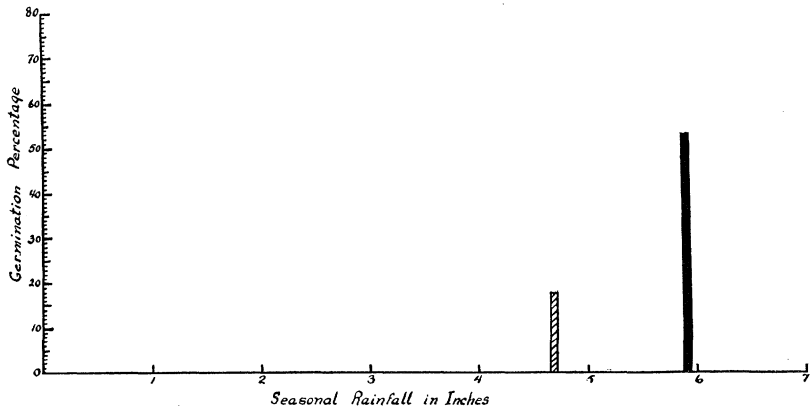
The areas allotted to the various stations are based on the records of the individual stations, supplemented by general observation throughout the year. Fig. 4 shows the approximate area covered by the different amounts of rainfall.



Comparison of Germination Percentages of
Hilaria mutica No. 1 and No. 2

Legend	<i>Hilaria mutica</i> No. 1 ▨	<i>Hilaria mutica</i> No. 2 ■
	Seasonal rainfall 3.88 inches	Seasonal rainfall 4.27 inches
	Germination 34%	Germination 91%

FIG. 1



Comparison of Germination Percentages of
Aristida longiseta No. 13 and No. 15

Legend	<i>Aristida longiseta</i> No. 13 ▨	<i>Aristida longiseta</i> No. 15 ■
	Seasonal rainfall 4.71 inches	Seasonal rainfall 5.94 inches
	Germination 17%	Germination 54%

FIG. 2

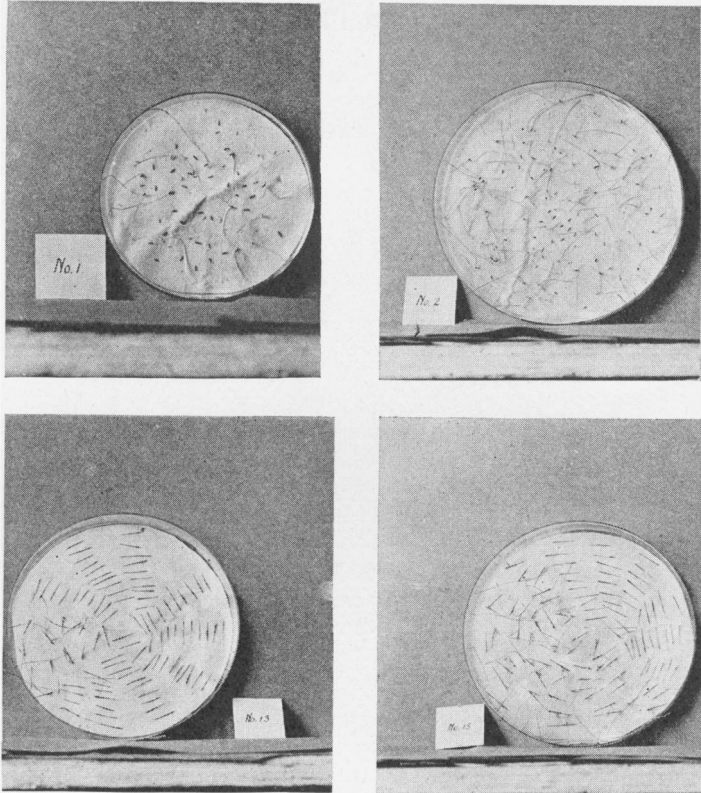


FIG. 3.—Upper row: comparison of germination of *Hilaria mutica* no. 1 (50 seeds, 34 per cent germination) and no. 2 (100 seeds, 91 per cent germination); lower row: comparison of germination of *Aristida longiseta* no. 13 (100 seeds, 17 per cent germination) and no. 15 (100 seeds, 54 per cent germination).

Seasonal Precipitation Map for 1926

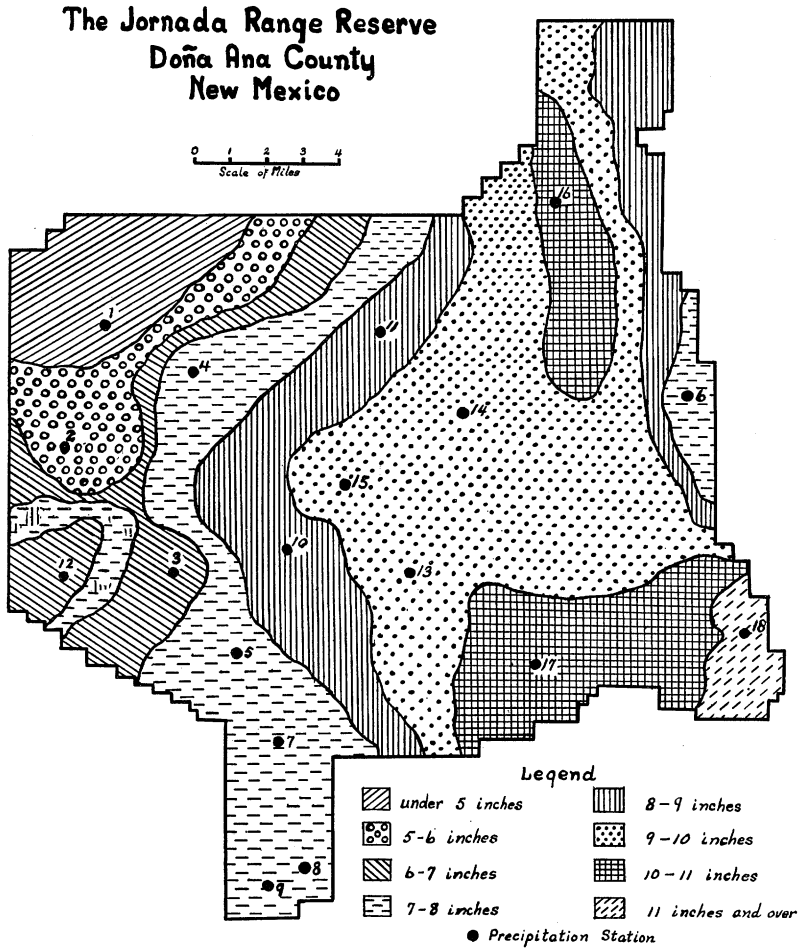


FIG. 4

Annual Precipitation Map for 1926

The Jornada Range Reserve
Doña Ana County
New Mexico

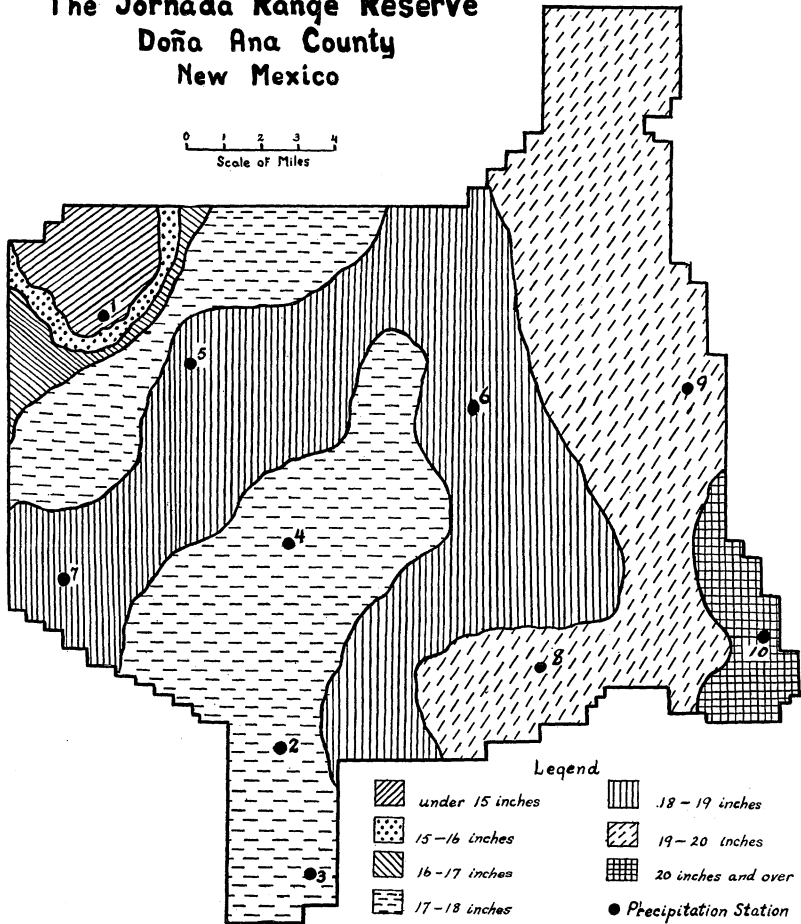


FIG. 5

The areas allotted to the various stations are based on the records of the individual stations, supplemented by general observation throughout the year. Fig. 5 shows the approximate areas covered by the different amounts of rainfall.

TABLE XXV
ANNUAL PRECIPITATION ON JORNADA RANGE RESERVE, 1926

No.	STATION	PRECIPITATION (INCHES)	RANGE (INCHES)
1.....	Red Lake	14.67	Under 15 15-16 16-17
2.....	South Well	17.30	17-18
3.....	Stuart Well	17.39	17-18
4.....	Headquarters	17.42	17-18
5.....	Middle Well	18.29	18-19
6.....	Road Tank	18.33	18-19
7.....	West Well	18.83	18-19
8.....	Ragged Tank	19.48	19-20
9.....	Ropes	19.68	19-20
10.....	St. Nicholas	21.17	Over 20

Discussion

From the tables of the various tests can be ascertained the germination percentages under different conditions. Table VII gives the final germination percentage for the seeds collected in 1926 and table IX for those gathered in 1925. From all the results it may be seen that the two lots of *Hilaria mutica* (nos. 1, 2) show a decided difference in their germination percentages. In no. 1 the germination percentage is always lower than in no. 2. By referring to the annual and seasonal rainfall figures, especially to the seasonal which includes just the rainfall during July, August, and September, when growth is taking place, one finds that no. 2 was collected in an area which received 2.69 inches more for the whole year than did the area in which no. 1 was growing. This probably accounts for the difference in the germination percentage, even though no. 2 was collected 19 days sooner than no. 1. The same is true of *Aristida longiseta* nos. 13 and 15. No. 15 was collected 16 days sooner than no. 13, but it gives much better germination results. The seeds of no. 15 received 1.23 inches more rainfall during the growing season than did the no. 13 seeds.

The *Bouteloua* (*B. eriopoda* and *B. gracilis*) had no seeds, so there was no germination percentage for either of them. Although tests were made in which the entire spikes were placed in sterilized Petri dishes in the 25° C. oven, no germination results were obtained. It is known that the *Bouteloua* grasses seed rarely, their means of propagation being vegetative rather than sexual.

Tables V and X show that germination is not increased by higher temperatures, as 35° C., nor does the alternation of low and high temperatures, that is from 5 to 25° C., increase germination (table XVIII). The best results are at 25° C.

In regard to the *Sporobolus* seeds nos. 9, 12, 14, 22, and 23, which have an unusually hard seed coat, it was found necessary to prick them in order to obtain germination results. Shaking the seeds in bottles containing coarse sand helped germination somewhat, but the writer believes that much better results could be obtained if the period of shaking were considerably extended. This method would scratch the coats of the seeds in a way similar to that in nature.

Summary

1. The amount of rainfall during the year, especially during the growing season and at the time of harvesting, affects the vitality of the seed.

2. *Bouteloua eriopoda* and *B. gracilis* do not seed very often. Most of the florets are sterile, and because of their similarity to the fertile florets, are hard to distinguish from them.

3. *Aristida* seeds germinate just as well in the light as they do in the dark.

4. The seed coat is important in *Sporobolus* seeds, as it keeps out water and prevents germination. The seed coat must be punctured by some means before good germination results. Soaking affects the seed coats but little; shaking even for 9 hours in sand has little effect; and scratching or pricking hastens germination greatly.

5. The seeds of *Sporobolus airoides* do not need pricking to produce good germination results. The seed coat is more permeable to water than are the seed coats of the other species.

6. The *Sporobolus* seeds from 1925 retained their vitality very well.

I wish to thank Mr. J. D. SCHOELLER, the Director of the Jornada Range Reserve Station, for supplying the seeds used in this investigation and for the climatic and geographic data of the region. I also wish to express my gratitude to Professor H. C. COWLES for his continued interest and suggestion throughout the work.

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LITERATURE CITED

1. ATWOOD, W. MCK., A physiological study of the germination of *Avena fatua*. BOT. GAZ. 57:386-414. 1914.
2. CHASE, AGNES, First book of grasses. New York: McMillan Co. 1922.
3. CROCKER, WM., Rôle of seed coats in delayed germination. BOT. GAZ. 43:265-291. 1906.
4. DAVIS, OPAL HART, Some cases of delayed germination. Master's thesis. 1925.
5. DUVEL, J. W. T., The vitality and germination of seeds. U.S. Dept. Agric. Bull. 58. 1904.
6. ECKERSON, SOPHIA, Physiological and chemical study of after-ripening. BOT. GAZ. 55:286-299. 1913.
7. EVANS, CLYTEE R., Effect of temperature on the germination of *Amaranthus retroflexus*. BOT. GAZ. 73:213-225. 1922.
8. FAWCETT, H. S., Viability of weed seeds under different conditions of treatment, and a study of their dormant periods. Proc. Soc. Acad. Sci. 15:25-45. 1908.
9. GRIFFITH, DAVID, Grama grasses. Contrib. U.S. Nat. Herb. 14:343-428. 1912.
10. HILMAN, F. H., Nevada and other weed seeds. Nevada Agric. Exp. Sta. Bull. 38. 1897.
11. HIMMEL, E. N., Longevity of seeds. Master's thesis. 1921.
12. HITCHCOCK, A. S., A text-book of grasses. New York: Macmillan Co. 1914.
13. PACK, DEAN A., After-ripening and germination of *Juniperus* seeds. BOT. GAZ. 71:32-60. 1921.
14. PAMMEL, L. H., and KING, CHARLOTTE M., Delayed germination. Proc. Ia. Acad. Sci. 17:20-30. 1910.
15. WOOTEN, E. O., and STANDLEY, P. C., Flora of New Mexico. Contrib. Nat. Herb. 19:1915.
16. U.S. Dept. Agric., Office of Exp. Stations. Rules and apparatus for seed testing. Circ. 34. 1906.