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# Alpine and subalpine wetland plant communities of the Uinta Mountains, Utah

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# The Great Basin Naturalist

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### ALPINE AND SUBALPINE WETLAND PLANT COMMUNITIES OF THE UINTA MOUNTAINS, UTAH

George M. Briggs' and James A. MacMahon-

ABSTRACT.— Seven wetland areas in the subalpine and alpine regions of the Uinta Mountains are described. Most areas have a mosaic vegetation structure, comprised of several patches that are usually sharply demarcated from one another. Each patch has a distinct species composition, usually with only one or two species. Water level is associ ated with some of the vegetation patterns found in these sites. Standing crop varied from 28 to 360 g/m<sup>2</sup>. Sites show considerable variation in standing crop that could not be explained by elevation differences between sites. Sites with water flowing over them have substantially greater standing crops than sites where water stagnates. Seasonal patterns in shoot density and standing crop indicate one late summer peak in standing crop and little recruitment of shoots over the summer. On stagnant sites, the average stem weight shows <sup>a</sup> strong relationship to stem density. This pattern did not appear to be caused by thinning mortality and did not follow a "3/2 power law" pattern.

A conspicuous aspect of many of the west em North American mountain ranges is their wetlands. The Uinta Mountains of northeastern Utah have an abundance of wetland areas, from the lower elevations (around streams in sagebrush slopes) to areas near springs at high elevations in the alpine zone. The majority of wetlands are found at the heads of glaciated valleys in a zone just above and below treeline. The vegetation in these areas is dominated by members of the Cyperaceae (sedges), as is much of the upland region in the alpine zone of the Uintas. Al though both wetland and upland regions are dominated by Carex spp., the structure of the two regions is markedly different (Briggs and MacMahon 1982). In this study we describe a variety of sedge-dominated wetlands, detail some of their structural attributes, and dis cuss some of the factors that we think are important in determining their patterns of occurrence.

#### STUDY SITES AND METHODS

All sites are in the Uinta Mountains of northeastern Utah (40°45'N, 110°-111°W), the largest east-west trending mountain range in North America. Bedrock throughout the upper elevations of the range is quartzite. The range was heavily glaciated in Pleisto cene time (Hansen 1975) and is dominated by numerous large U-shaped valleys ending abruptly in steep-walled cirques.

No attempt was made to describe or cata log all the wetland areas of the Uintas. In stead, we chose to study a variety of wetland regions that varied in their species composition and standing crop. Most of these regions had a patchwork appearance because they were composed of regions (2-100 m<sup>2</sup>) of distinct species composition, often monotypic, which abruptly gave way to regions of a dif ferent species mixture. In sampling these areas, we first made a rough map that demarcated the "patches" found in them. Each

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patch was then sampled by the methods outlined below. We refer to each patch as a site and present figures representing the species composition and standing crop of each of these. The larger units of vegetation (the entire meadow composed of several patches) are not quantitatively described in this paper. Each site was given a two-letter code that represented the area from which it came and a number that indicated the specific patch sampled  $(e.g., SC-1)$  is patch 1 in area  $SC$ ).

The boundaries to each site were subjectively demarcated and a point was picked within the stand. A  $20 \times 50$  cm frame was placed at that point and all vegetation (ex cept bryophytes) was clipped at ground level. Nine additional samples were taken at regular intervals from the first point. These samples were sorted to species in the field, re turned to the lab, air dried for at least two months, and oven dried at 40 C for at least 24 h. Sorting to species was quite easy, even when based on vegetative characters, since there were rarely more than two species per sample. For larger species, the number of stems was directly counted and the average weight per stem calculated. For the smaller species, the average weight per stem was de termined by weighing three replicates of 100 stems. The total number of stems was obtained by dividing the total weight of that species in the sample by the weight/ stem. All sites were sampled in August, when the fruits of the dominant species were mature. Seasonal changes in aboveground standing crop, number of shoots, and number of fruiting shoots were determined in three of the sites by sampling three or four times during the summer. In addition, several individual plants were observed. The length of each leaf and the length and condition of the fruiting culm (if present) were measured at each sampling time.

At each site, soil samples were dug and pH of the soil was determined. All these sites were submerged at least part of the summer, and notes were made of both the length of time the site was submerged and the degree of water movement over the site.

#### Site Descriptions

Seven wetland areas were mapped and 21 sites were sampled within these areas. Each wetland area contains between one and five sites. Table <sup>1</sup> lists the dominant species, ele vation, and standing crop of all 21 sites. Fig ure <sup>1</sup> maps the vegetational patterns found in the more complex areas.

The highest wetland found was at an ele vation of 3,768 m, in an area adjacent to the steep, rocky talus pile that makes up the northeast side of King's Peak in the east central section of the Uinta range. Springs moist en the site all summer and feed a small stream that traverses the site. Large rocks are scattered throughout. The wetlands of this re gion consist of a series of small irregularly shaped pools containing 1-10 cm of water at the end of the summer. The emergent vegetation consists solely of Eriophorum scheuchzeri Hoppe. These pools were sampled as site ER-1.

To the north of King's Peak lies Henry's Fork Basin. The head of this basin has large expanses of sedge meadows and willow thickets interrupted by upland regions. In spite of a homogeneity in species composition (much of the area is dominated by Carex aquatilis), there is considerable variation in stature and standing crop of wet meadows within upper Henry's Fork Basin. We sampled two sites that represent extremes in Carex aquatilis meadows. HF-1 is <sup>a</sup> border to <sup>a</sup> small (20 m diameter) pond at an elevation of 3,278 m. Vegetation consisted of robust individuals of Carex aquatilis and Caltha leptosepala. Water stands at least 10 cm deep throughout the year, draining off the site to the north. HF-2 is a very different Carex aquatilis region, with a more dense stand of much smaller plants. There is no proximate open water and the ground surface was dry at the end of the summer although the ground was saturated with water at a depth of 2-5 cm.

The remaining five areas are within four miles of Mirror Lake on the west end of the Uintas. Area TL is part of <sup>a</sup> circular meadow approximately <sup>200</sup> m in diameter. The mead ow has a small stream meandering through itand has numerous small "oxbow lakes" and "kettleholes." The vegetation displays a pat tern commonly found in moist subalpine meadows of the Uintas: monotypic stands of Carex aquatilis and Eleocharis pauciflora, areas of Carex aquatilis and Eleocharis pauciflora mixed, and sections of bare ground, all

surrounded by an upland that has soil with less organic matter and a vegetation of Carex illota L. H. Bailey, Deschampsia cespitosa, and Ligusticum filicinum var. tenuifolium (S. Wats.) Mathias & Constance. A large portion of this area has standing water year around. This area was sampled in five sites (Fig. 1).

Areas FT and SE are small meadows in Picea engelmannii-dominated forests. The

vegetation in them is similar to that in TL and is mapped in Figure 1. Both areas were sampled as four sites. Area SC is a meadow in a Picea engelmannii-Pinus contorta-dominated forest. The meadow has a stream flowing through it and is kept moist by two large seepage areas. The vegetation presents a complex patterning, with part of the area ex hibiting the mix described at the TL site.

TABLE 1. Elevation, standing crop, and species found on the sites.



Other parts of the meadow have a dense turf contains a mix of *Carex illota* and *C. aqua*-<br>of *Scirpus cespitosus* L. associated with *Carex* tilis. The whole meadow has standing water<br>*aquatilis, Eleocharis pauciflora*,



mollis, CAAQ = Carex aquatilis, CAIL = C Fig. 1. Maps of four of the wetland areas: A, Area SE. B, Area TL. C, Area SC. D, Area FT. ARMO = Arnica  $t$ osa, ELPA = Eleocharis pauciflora, JUPA = Juncus parryi, POPA = Poa pratensis, SCCE = Scirpus cespitosa.

Area AQ is on the margin of <sup>a</sup> small pond that narrows into a stream at one end. It is surrounded by a Picea engelmannii-Pinus contorta-dominated forest. This area is at least 30 cm under water in the spring. Standing water is present to a depth of 10 cm in the fall. One site was sampled in this area, a region of nearly pure Carex aquatilis.

#### Results and Discussion

#### Vegetation

The number of species sampled on each of the 21 sites (Table 1) varied from one to seven. Most of the species present in the wetland sites are common species of the western United States. The Carex species collected can be separated, phytogeographically, into two groups: western cordilleran species {Carex nigricans, C. illota) and circumboreal species  $(C.$  rostrata,  $C.$  aquatilis, and  $C.$ canescens).

One of the striking features of these wetlands is the dominance by plants that are ca pable of extensive spread by rhizomes. The mosaic structure (Fig. 1) of these communities is a function of the strongly rhizomatous nature of the dominant plants and some, perhaps most, of the "patches" seen represent single plants (genets. Harper 1977) formed by growth from a single propagule. The patterns seen in these wetlands are probably dependent on both the history of propagule arrival and on variations in environmental conditions affecting the success of particular species. A complicating factor results from the "inertia" of these sites—a re sistance to vegetational change resulting from both the harsh nature of the climate and the vigorous mode of wetland plant growth. Both these factors could make established vegetation patterns difficult to disrupt. Al though seedling establishment is common in some wetlands (Lieffers and Shay 1982), both Costello (1936) and Bernard (1975) noted that seedling establishment was rare in some Carex wetlands. Our sites included areas where the dominant was in very poor condition (e.g., HF-2) or had died off completely, leaving areas of bare ground (e.g., part of TL). Both biotic and abiotic factors at these sites may change with time. These changes

will sometimes be to the detriment of the species that has been dominant. But, because of the difficulty in establishing additional species, the dominant may continue to exist in <sup>a</sup> depauperate condition and may even die back completely before another species is able to invade the area. Thus, the species present today may not reflect present conditions but may represent conditions of an earlier time.

One environmental variable that we found associated with community structure is water level. In TL and FT the Carex illota section was always upland relative to the Carex aquatilis-Eleocharis pauciflora sections. In SC the part of the plot containing Scirpus cespitosus was on higher ground than those regions containing C. aquatilis and Eleocharis pauciflora. Some species distributions could not be associated consistently with water level. Carex aquatilis could be found in areas with both more and less standing water than the often adjacent E. pauciflora areas (areas TL, FT). Carex illota, although usually restricted to areas with no standing water (FT-4, TL-2), sometimes was found in areas 5 cm deep in running water (SC-4).

It is generally thought that the influence of water level in wetland regions is based on variations in waterlogging and aeration. Soil aeration may influence plants either by af fecting root respiration (Mendelssohn et al. 1981), which could affect water and nutrient flow through roots, or by affecting soil nutri ent status. The influence on soil nutrition can occur directly, by influencing the chemical forms available and their solubilities (Jones 1971, 1972), or indirectly, by affecting soil microbial activity.

The importance of aeration on species dis tribution is represented in our observations on Carex illota, which is intolerant of waterlogged soils if the water is stagnant (and pre sumably poorly aerated), yet grows quite well under 5 cm of water when it is flowing over the site (and probably better aerated). Further evidence for the importance of aeration in these sites is shown in standing crop data presented below. If aeration does in fluence the structure of these communities, the complex nature of some of these sub alpine sites could be due to underground water flow patterns and resultant aeration patterns. These flow patterns may be due to



#### MOISTURE CLASS

Fig. 2. Graph showing the relationship between standing crop on <sup>a</sup> site and <sup>a</sup> measure of water movement over that site.

springs and may not be readily discemable without measurement of soil conditions.

The peak aboveground standing crop val ues (Table 1) varied considerably, both between areas and also between adjacent sites in one area (e.g., area SC). The very low standing crop value for site ER was undoubtedly due to the short growing season and low temperatures associated with this, the highest site. However, aside from site ER, there is very little relationship between elevation and peak aboveground standing crop. Gorham (1974) derived a regression equation relating standing crop of *Carex* meadows to the highest monthly mean temperature of the year. His data on 11 pure Carex stands (on both rich and poor soils) fit the regression line well  $(r = 0.84)$ . The data of Auclair et al. (1976) on wetland stands that were not pure Carex also fit this regression. Using weather data for areas in Colorado that are comparable to ours, Gorham's equation predicts a standing

crop of  $292 \text{ g/m}^2$  for a site at 3,109 m  $(10,200 \text{ ft})$  and 235 g/m<sup>2</sup> for a site at an elevation of  $3,566$  m  $(11,200)$  ft). The standing crop values that we found are both above and below these values, with most sites below. Gorham's equation is most appropriate for those sites where water is not stagnant (HF-1, SE-2, SE-3, SC-4). The sites with less water movement have less standing crop than predicted by Gorham. The reduced growth on sites where water stagnates is shown in Figure 2, where standing crop is plotted against a gradient in the amount of water movement through the site. Reduced growth on the stagnant sites is probably due to low oxygen levels that may be disturbing root res piration, mineral uptake, or soil nutrient status.

An oxygen-limited situation in Carex meadows might explain the timing of nutri ent uptake in wetlands. Boyd (1970) and Ber nard and Solsky (1977) note considerable



Fig. 3. Seasonal trends in aboveground standing crop for two of the pure Carex aquatilis sites, SE-2 and SE-3.

mineral uptake by wetland plants in the early spring. Boyd hypothesized that this early mineral uptake (preceding the growth period) was an adaptation to allow certain species to procure nutrients before other species. Another possible reason for mineral uptake early in the spring could be related to oxygen concentrations. In the early spring, oxygen in wetland soils may be at its highest levels be cause of well-oxygenated runoff waters and because low temperatures increase the amount of oxygen that water can hold. Thus, plants may take up nutrients in the early spring because that is the most favorable time for root respiration.

In several of the Carex aquatilis-dominated sites, we observed the variation in standing crop during the summer. The typical pattern in these subalpine meadows (Fig. 3) includes a single peak in aboveground standing crop which occurs in late summer. Gorham and Somers (1973) and Bernard and MacDonald (1974) also found such a pattern for wetlands composed of C. aquatilis and C. lacustris, respectively. On our sites there was relatively little new shoot production over the course of the summer (Fig. 4), and the changes in aboveground standing crop were being caused by the growth of individual shoots. Some Carex species (e.g., C. rostrata) have biannual periods of shoot production and



Fig, 4. Seasonal trends in stem density (stems/m^) for Carex oquatilis on SE-2 and SE-3.

these may be responsible for bimodal patterns in biomass (Gorham and Somers 1973), although other factors may cause bimodal patterns. The pattern in shoot production that we observed for C. aquatilis had been found previously for this species (Gorham and Somers 1973), as well as for C. lacustris (Bernard and MacDonald 1974), and would be expected in regions with short growing seasons such as the one we studied.

Based on our observations, the life history of C. aquatilis is initiated by growth of a new shoot sometime during the winter or spring. The shoots grow throughout the summer and then overwinter. The following spring the old shoots are distinguishable from shoots of the current year by their large size and old leaves. One or two of these old leaves may undergo additional growth in the spring. At least some and possibly all of the shoots un dergo two or possibly more seasons of growth before they flower or die. Many shoots may not flower. Floral initiation is evident very early in the spring; floral parts are visible less than two weeks after growth initiation. During the growing season leaves are initiated, grow quickly to a maximum length, and re main green until early September.

There was considerable variation in plant density in the C. aquatilis stands. On the stagnant sites (those where water was not flowing), the variation in density was related to stem weight. If a plot is made of the natural logarithm of plant density vs. the natural logarithm of plant weight (Fig. 5), it can be seen that the stagnant sites fall on a line



Fig. 5. Graph showing the relationship between stem density and average stem weight for five Carex aquatilis stands where water was stagnant.

 $(r^2 = 0.98)$ . Sites with flowing water do not fall on this line and have substantially greater plant weights at a particular density than those on stagnant sites. The regression line of Figure 5 describes the following equation:

[1] 
$$
w = 5.53\rho^{-0.86}
$$
  
where  $w =$  individual plant  
weight;  $\rho =$  plant density

The form of this equation is characteristic of monotypic stands and an exponential value of -3/2 has been found for a wide variety of species of differing life forms and habitats (Gorham 1979, White 1981). The equation has been labeled the "3/2's thinning law" be cause it describes the changes in density and plant weight that occur in stands during self thinning. On our plots it is likely that the pattern of density and plant weight is not being caused by mortality (thinning) but rather by recruitment (see Fig. 4). Gorham (1979) cites studies with imcrowded conditions, and therefore little self-thinning, where the exponent value in equation (1) is closer to -1. Our stagnant sites appear to be similar areas. With an exponent of  $-1$ , standing crop (w.  $\rho$ ) remains constant regardless of density. Such sites may represent areas where stand ing crop is limited by factors (such as fertility?) that prevent densities to reach levels

where self-thinning in the typical 3/2's fash ion can occur. The precise factors that dictate the 3/2's thinning law are still not eluci dated (White 1981). Perhaps studies on such areas as these, where the "law" does not hold, may help to clear up the problem.

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