





quinoa have highly nutritious characteristics [23]. Quinoa is a dicot plant, and its tallness is 1.5-6.5 ft. Its cropping season is from 15 Oct-30 March. The temperature range for this crop to grow is 12-20°C and its pH is equal to 8. It is cultivated in northern areas of Pakistan like Hunza and small plain areas like Bahawalpur. Quinoa leaves are broad, and these generate edible seeds and leaves [8]. The protein content in quinoa is about 12.5-16.7% and the fat content is about 5.5-8.5% [24].

Quinoa yield is about 2.5 tonnes per hectare in Pakistan. A single spikelet yields 200 g of quinoa grain [25]. It contains 71% water, 21% carbohydrate (Starch form), 4.4% protein, and 1.6% unsaturated fatty acids [26]. Grains of quinoa are rich in omega-3 and omega-6 fatty acids. Quinoa has 120 color varieties but red, black, and white are mostly used [27]. Quinoa is a cereal crop, and it can withstand harsh climatic stress conditions. It can tolerate the worst temperatures and lower temperatures such as from -40°C to 350°C. Quinoa is grown in sandy to loamy sand-type soil [28]. It is a  $C_3$  halophytic type plant. It is rich in protein content and amino acids.

The present research was primarily based on the theory that Si and SA can be utilized to reduce the adverse impacts of water shortage conditions on quinoa, thus increasing its production. By reducing the negative impacts of water shortage conditions in quinoa using Si and SA treatment, the current work filled the research gap. Researchers have done experiments for determining Si and SA impacts in reducing water deficit stress in other crops but their interactive effect in quinoa is not widely studied. Furthermore, work on comparative studies to evaluate the effectiveness of silicon and salicylic acid for reducing water deficit stress in quinoa is scarce. Based on this discussion, the present study was conducted to determine the impact of the combined application of silicon and salicylic acid for improving quinoa production, grain output, and development under water shortage stress.

## Experimental Procedures

### Experiment Setup

A pot experiment was carried out in the wirehouse of ISES, UAF. Each container was filled using 0.7 kg of soil and two water levels (100 and 60% FC) were maintained in pots by using a weighing method to develop water deficit conditions in the soil. Quinoa (*Chenopodium quinoa*) variety “V-17” was used. The suggested dose of NPK fertilizers for quinoa (i.e., 75-50-50 kg ha<sup>-1</sup>) was applied to all treatments before sowing. Irrigation was done in measured quantity to the pots according to the moisture condition of the soil. Three levels of silicon i.e., control, 100 and 200 mg kg<sup>-1</sup> using potassium silicate as a source were used. Salicylic acid (1 mM) was foliar applied at the vegetative stage. Three foliar sprays with

an interval of 10 days were done. There were thirty-six pots arranged in a completely randomized design (CDR) with factorial arrangements comprising 12 treatments in triplicate.

### Measurement of Physiological Parameters

At the end of the vegetative stage, physical parameters such as chlorophyll (SPAD Value), relative water (RWC), and membrane stability (MSI) were investigated. Total chlorophyll contents (SPAD Value) in quinoa leaves was measured at the completion vegetative growth stage by the Soil Plant Analysis Development meter 502, which works on the light emission and transmission principle. For the determination of chlorophyll contents, three mature leaves of the quinoa plant were selected and placed under the SPAD meter. The average SPAD measurement of the leaf was determined [29]. Relative water content (RWC) was measured after the vegetative stage. For the measurement of RWC young leaves from the plant were taken and measured, following the procedure given by Barrs and Weatherly [30]. To calculate MSI, the method used was given by Sairam et al. [31].

### Measurement of Growth Parameters

Crop was harvested as it reached maturation. Crop growth responses such as fresh and dry weight of shoot and root, plant height, and length of root were determined at the time of harvesting. Initially, samples were air-dried and then oven-dried for 72 h to determine shoot and root dry weight.

### Measurement of Yield Parameters

Crop yield responses such as panicle length and grain yield were determined. Panicle length was calculated at harvesting time while grain yield was determined after oven-dried samples.

### Determination of Na<sup>+</sup> and K<sup>+</sup> in the Shoot

For the determination of Na<sup>+</sup> and K<sup>+</sup> in shoot wet digestion was required which was done according to the method of Gargari et al. [32]. Na<sup>+</sup> and K<sup>+</sup> concentration was evaluated by using a Sherwood-410 flame photometer [33].

### Statistical Analysis

Statistix 8.1 software was used to statistically examine data. The overall significance of data was assessed using the analysis of variance (ANOVA) method, and LSD was applied to compare the results at a 5% level of significance. All the analyses of various parameters were conducted using three replicates. The  $p$ -value<0.05 shows that there were significant differences among treatments [34].





















