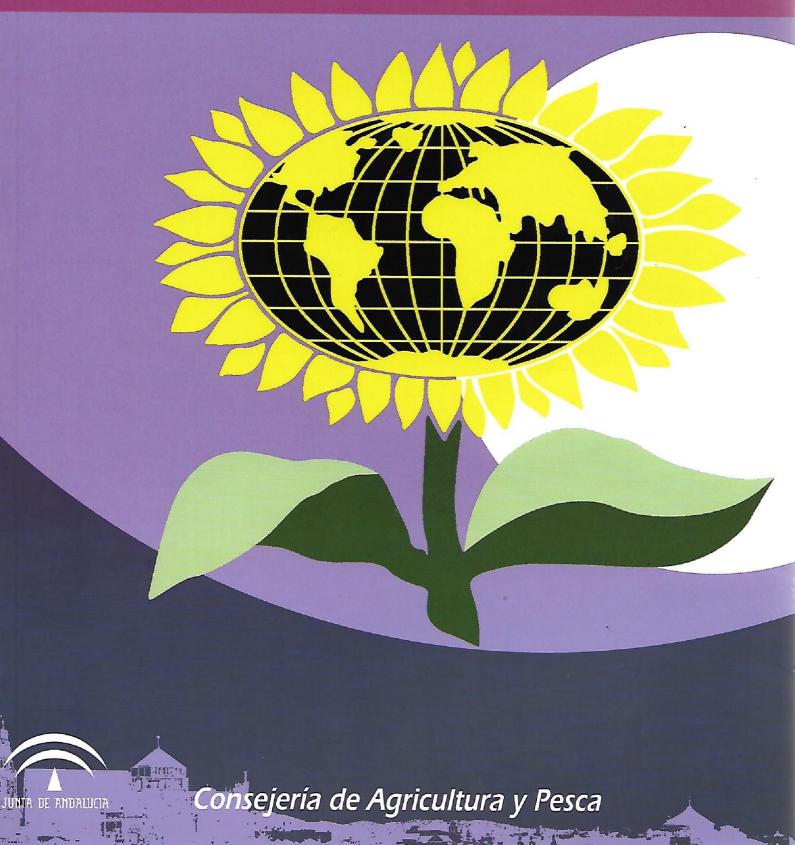
CULTIVOS HERBACEOS

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Vol. 1



Proceedings of the 17th International Sunflower Conference

Vol. 1



Córdoba, Spain June 8-12, 2008

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Instituto de Investigación y Formación Agraria y Pesquera CONSEJERÍA DE AGRICULTURA Y PESCA





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Photography: Spanish landrace of confectionary sunflower collected by L. Velasco and B. Pérez-Vich in Villarta de San Juan, Ciudad Real, Spain, on October 10, 2007.

Foreword

The proceedings of the 17th International Sunflower Conference contain 142 contributions from scientists of 24 countries. They include plenary lectures in several disciplines and regular communications presented in posters during the conference and discussed in the corresponding workshops. The manuscripts are classified by disciplines. They offer a good picture of the current state of the art of sunflower research and cultivation around the world.

The manuscripts in the *Proceedings* have been reviewed by an editorial committee with the main objective of helping the authors to improve their manuscripts through a critical reading. The authors received the edited manuscripts together with the comments of the reviewers and then went on to draft their final version. All the manuscripts received have been published in the *Proceedings*. The contents of the manuscripts are the responsibility of the authors. They should be considered as being privileged communications that require the express consent of the authors to be reprinted in part or as a whole. We wish to thank both the members of the Editorial Committee for their dedication to the task of editing such a large number of manuscripts, as well as all the authors for their collaboration throughout the whole edition process.

The Organizing Committee would also like to thank Diana Badder and José A. Palacios for their excellent editorial assistance in the preparation of these *Proceedings*. We are indebted to the Spanish Association of Sunflower Breeders (Asociación Española de Mejoradores de Girasol), which collaborated actively in the organization of the conference, and, very especially, to Juan Parejo, who was in charge of the financial side.

Finally, we would like to thank all the participants in the conference, who have contributed to its success by a careful preparation and revision of manuscripts and posters, presentation of their research in the workshops, and stimulating discussions throughout the conference on the scientific and technical aspects of sunflower research and cultivation in the world.

The Organizing Committee 17th International Sunflower Conference Córdoba, Spain. June 8-12, 2008

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The appropriate technique for collecting and measuring the amount of floral nectar in sunflower (*Helianthus annuus* L.)

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ABSTRACT

The available techniques for collecting and measuring the amount of floral nectar are applicable but often found to be unrepresentative. Centrifugation yields larger samples but they also include nectar that is not actually accessible to insects, the capillary method has been described as unsuitable because of possible damage to the nectary tissue, the method including filter paper is considered to be unreliable because of evaporation and nectar extraction methods including washing are considered limited because the solution may include sugars from plant tissue cells. We have found that capillary tubes with inner diameter of 0.25-0.5mm and outer diameter of 0.5-0.75mm are suitable for nectar collection in sunflower. To determine the amount of nectar, we isolate five inflorescences per sunflower line at the start of flowering and collect the nectar two days after the isolation. The capillary tube is inserted between the style and filaments down to the nectary. After the level of nectar stops rising the next flower is processed. The tubes can be measured on an analytical scale and the amount of nectar is obtained as the weight increase in comparison to the empty tube. Faster determination of floral nectar amount can be provided by using calibrated capillary tubes of a known and uniform inner diameter. The appropriate outer diameter of the capillary tubes reduces the risk of tissue damage and allows more precise collecting so that the capillary method is preferable to others for nectar collecting in sunflower.

Key words: capillary technique - nectar quantity - sunflower

INTRODUCTION

Sunflower is one of the plant species that produces pollen which is too heavy for wind dispersal (Putt, 1940). Even though the cultivated sunflower has a reasonable percentage of self-compatibility it still benefits from insect pollination. One of the major components influencing pollinator choice is certainly the production of nectar, whose amount and quality are often studied.

The nectaries in *Asteraceae* family form on top of the ovary and surround the style base (Mani and Saravanan, 1999), (Fig. 1). The nectar can be accessed for quantification purposes by capillary tubes (Hocking, 1953), volumetric centrifugation (Bosi, 1973), filter paper strips (McKenna and Thomson, 1988), flushing of water into the corolla (Cresswell and Galen, 1991) or floating the flowers inverted in water (Manetas, 2000) depending on flower structure. These techniques were used with a variable success, but the overall conclusion is that no single method can be considered satisfactory for all plant species (Mesquida et all, 1988). The capillary method can be used in sunflower but it is necessary to use capillary tubes of appropriate dimensions. If the outer diameter is too large it is not possible to access the nectar without destroying the surrounding corolla tissue (Fig. 1.) and if the tube is too thin then the intake of nectar is slower and the strong capillary force may make the extraction of nectar from the tubes difficult.

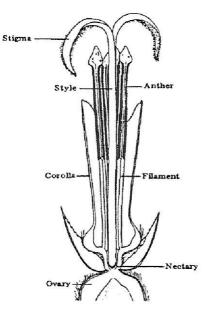


Fig. 1. Longitudinal section of a sunflower disk flower showing the location of the nectary

MATERIALS AND METHODS

We found that capillary tubes with inner diameter of 0.25-0.5mm and outer diameter of 0.5-0.75mm are suitable for nectar collection. The length of the tube then determines the capacity and can be picked for its suitability to the collecting design but should not be smaller than 15 μ l.

To determine the amount of nectar, we isolate five inflorescences per sunflower line at the start of flowering with linen bags to prevent insects from collecting nectar. The best moment is when the first two rows of disk flowers have opened. Two days after the isolation, at approximately 8 AM, the inflorescences are cut and taken to the laboratory in a portable refrigerator to minimize evaporation and the change in nectar volume. It is advisable for the transport duration to be as short as possible. The following should be prepared to analyze one inflorescence:

- 1. Four capillary tubes (previously weighed on an analytical scale) each placed in a separate tube labeled with sample and replication to ease the work of collecting and measuring
- 2. A clean vegeglass (previously weighed on an analytical scale)
- 3. A clean HPLC vial with sample label on it filled with 1 ml mixture of AcCN:H₂O in a ratio of 75:25
- 4. A plastic dish with sample label on it for deep freezing

Four groups of five analyzed flowers are equally far from each other on an inflorescence. We collect nectar from 5 fully opened disk flowers with one non calibrated capillary tube. The tube is inserted between the style and filaments down to the nectary (Fig. 1.). After the level of nectar stops rising the next flower is processed. When a total of 20 flowers are finished, the tubes can be measured on an analytical scale and the amount of nectar is obtained as the weight increase in comparison to the empty tube. Faster determination of nectar amount in flowers can be provided by using calibrated tubes (capillary tubes with a uniform known inner diameter) for nectar collecting, in which case the height of nectar in tubes can be correlated with the nectar volume. This method is suitable when it is necessary to determine the amount of nectar in field conditions, without cutting the sunflower head and taking it to the lab.

The next step in method developing is a qualitative and quantitative HPLC analysis of nectar extracted from a single inflorescence as a collection from 20 disk flowers. For this purpose, ten disk flowers are also pulled off with tweezers, put in a glass and weighed on an analytical scale to obtain the information about the flower mass and possible correlation with nectar production.

The nectar collected in capillary tubes, after weight measurement, can also be kept for subsequent analysis. The contents of all capillary tubes from a single inflorescence are transferred into a HPLC vial (2 ml) filled with 1 ml mixture of $AcCN:H_2O$ in a ratio of 75:25 and placed in a refrigerator for

subsequent HPLC nectar quality analysis. Twenty flowers are pulled out of the disk with tweezers, frozen in liquid nitrogen and then kept at -72°C.

DISCUSSION

The rest of the techniques cited are applicable but often unrepresentative. Centrifugation yields larger quantities but they also include nectar that is not actually accessible to insects and modified chemical composition due to tissue lesion (Mesquida et al., 1988). The method including filter paper is considered to be unreliable because of evaporation (Livtzieva, 1954). Nectar extraction methods including washing are considered limited because the solution may include sugars from plant tissue cells (Kenoyer, 1917).

A combination of capillary method and filter paper can be used so that the nectar is extracted with capillary tubes and than ejected on to a filter paper, which is measured for total nectar. After the evaporation has finished the amount of sugars is obtained as a difference between wet and dry filter paper.

The capillary method has been used on sunflower (Pham-Delegue et al., 1985) but it has also been described as unsuitable for collecting nectar amounts less than 1 μ l and to cause damage to the nectary tissue (McKenna, 1988). The appropriate outer diameter of the capillary tubes reduces the risk of tissue damage and allows more precise collecting so that the capillary method is preferable to others for sunflower nectar collecting.

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