## PROPAGATION OF CAPER BY SEED, CUTTINGS, OR BY TISSUE CULTURE\*

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## ABSTRACT

Caper seeds were immersed first in H<sub>2</sub>SO<sub>4</sub> and KNO<sub>3</sub> and then in GA<sub>3</sub> or in the GA<sub>3</sub> alone for various times and caper cuttings were treated with increasing doses Indolacetic acid (IAA), Indolebutyric acid (IBA) and 2-Naphthelene acetic acid (NAA-). The cuttings and the seeds were grown under *in vivo* and *in vitro* conditions. The highest seed germination percentage (78.6%) was obtained from the treatment with GA<sub>3</sub> (500 ppm) for 24 hours. The cuttings treated with NAA for 7 minutes resulted in 51.6% rooting in perlite. The IBA treatment (6 ppm) applied to 1-1.5 cm explants gave 53.8% rooting in MS medium. Development of the 96 plants grown in the experimental field was found to be at high level.

Key words: Capparis spinosa L., caper development, rooting