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25 **Use of 1-Methylcyclopropene in cyclodextrin-based nanosponges to control grey**
26 **mould caused by *Botrytis cinerea* on *Dianthus caryophyllus* cut flowers**

27

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43

44 **Abstract**

45 *Botrytis cinerea* is one of the pathogens resulting in the heaviest commercial losses in
46 ornamental cut flowers, and the severity of grey mould disease partly depends on the
47 presence of ethylene in the storage environment. The efficacy of β -cyclodextrin-based
48 nanosponge 1:8 (NS) - 1-methylcyclopropene (1-MCP) complex was evaluated as a
49 novel control agent in protecting carnation (*Dianthus caryophyllus* L. 'Idra di

50 Muraglia') cut flowers against *B. cinerea* infection. Two concentrations of this
51 innovative, non-volatile 1-MCP formulation (NS complex, 0.25 and 0.5 $\mu\text{L L}^{-1}$, a. i.)
52 were compared to the commercial gaseous 1-MCP (0.25 $\mu\text{L L}^{-1}$, a. i.), and to an
53 inoculated control. Furthermore, a not-inoculated control was used to assess the natural
54 infection level. Eleven days after inoculation, the development of grey mould on
55 carnation was significantly reduced (59.9% of flower surface) in cut stems treated with
56 the NS complex at low dosage, compared to the high dosage of the NS complex
57 (91.5%), the commercial gaseous 1-MCP formulation (76.2%) and to the inoculated
58 control (100.0%). Endogenous ethylene production was correlated to the symptoms
59 development. Results showed a reduced ethylene production in 1-MCP treated flowers
60 (0.25 $\mu\text{L L}^{-1}$, a. i., both suspended and gaseous formulation). NS complex could
61 therefore be an effective alternative to conventional chemicals to protect ornamental cut
62 flowers.

63

64 *Keywords:* carnation, grey mould, nanocarriers, postharvest, ethylene antagonist.

65

66 *Abbreviations:* CD, cyclodextrin; 1-MCP, 1-methylcyclopropene; CD-NS, β -CD-based
67 nanosponge 1:8; CD-NS complex, β -CD-based NS 1:8 - 1-MCP complex

68

69 **1. Introduction**

70 The flower trade worldwide is marked by an increasing competitiveness in cut flower
71 offer (Serra, 2003). An important cut-flower crops grown internationally is carnation
72 (*Dianthus caryophyllus* L.). Increasing attention is paid to postharvest vase life, that
73 plays a crucial role in the quality of cut flowers. The senescence process is induced by

74 several factors, among these are ethylene and pathogens (Woltering and van Doorn,
75 1988; Serek et al., 1995a,b).

76 *Botrytis cinerea* is an airborne pathogen for a wide variety of cut flower crops,
77 causing grey mould responsible of severe economic losses. During the last few decades,
78 the introduction of new ornamentals, and the improved growing techniques caused
79 significant changes that contributed to aggravate the severity of diseases (Daughtrey and
80 Benson, 2005). Therefore, a high priority on the research of sound management of
81 greenhouse- and nursery-grown ornamentals, and of a more effective disease control, is
82 required.

83 Considering the increasing restrictions on the use of pesticides (European Regulation
84 1107/2009 and Directive 2009/128, US Food Quality Protection Act), the development
85 of new eco-friendly disease management strategies is needed. In a previous study, the
86 use of the ethylene antagonist 1-methylcyclopropene (1-MCP) resulted effective in
87 reducing damages caused by *B. cinerea* in cut flowers of several ornamental species
88 (Seglie et al., 2009). However, difficulties in the application of this gaseous compound,
89 such as the necessity of enclosed areas to prevent gas leakage, the need of continuous or
90 repeated treatments, and the low action at temperatures (0–5 °C), complicate its
91 commercial use (Serek and Sisler, 2005; Serek et al., 2006).

92 The inclusion of 1-MCP in cyclodextrin-based nanosponges structures (CD-NS,
93 patented by Trotta et al. ultrasound-assisted synthesis of cyclodextrin-based
94 nanosponges patent WO2006/002814) can reduce these practical limitations, and
95 already showed to be effective in prolonging cut flower vase life (Seglie et al., 2011a).
96 CD-NS is a delivery system able to induce an extended release of the 1-MCP, leading to
97 benefits such as reduced active ingredient dosages required and reduced number of
98 delivery times as compared to the gaseous commercial product.

99 In the present study, the effectiveness of the non-volatile formulation of 1-MCP
100 included in CD-NS (CD-NS complex) in controlling *B. cinerea* damage on carnation cut
101 flowers was evaluated.

102

103 **2. Materials and methods**

104 For the experiments, carnations (*Dianthus caryophyllus* L. 'Idra di Muraglia') were
105 grown in standard greenhouse conditions in Sanremo, Liguria, Italy. Cut flowers were
106 harvested at maturity stage (vertical sepals, vivid petal and stem colour) and taken to the
107 postharvest laboratory within 24 h, where they were re-cut and labelled. The experiment
108 was performed three times, each treatment included 3 repetitions of six cut flowers
109 (stems 30 cm long). Twelve carnation cut flowers, inoculated or not with the pathogen,
110 were kept in tap water as controls.

111 Stems were then placed in vases with a suspension of 1-methylcyclopropene (1-
112 MCP) included in β -cyclodextrin-based nanosponges 1:8 (CD-NS complex, 6% a. i.) at
113 two different concentrations of active ingredient (0.25 and 0.5 $\mu\text{L L}^{-1}$, a. i.), or exposed
114 to 6 h treatment with the commercial gaseous 1-MCP (3.3% a. i., SmartFreshTM,
115 AgroFresh Inc., USA). Treated flowers were inoculated with a *B. cinerea* conidial
116 suspension (10^4 conidia mL^{-1}) to favour the mould development.

117 Daily, the extent of *B. cinerea* development on each flower was monitored, counting
118 the number of infected petals in relation to a mean total number of petals (61.9),
119 calculated on 10 flowers.

120 Ethylene production was daily measured by keeping single treated flowers in air tight
121 vases (250 mL) containing 50 mL of the different preservative solutions, or tap water
122 for the controls. The ethylene concentration was measured using a digital Agilent
123 Technologies gas chromatograph, 6890N Network GC system (Santa Clara, California).

124 The gas carrier was N₂ at 40 mL min⁻¹, and the column temperature was 60 °C. For
125 each treatment, three samples were considered.

126 Statistical significance among mean values was assessed performing the analysis of
127 variance (ANOVA), and the Ryan-Einot-Gabriel-Welsch's multiple stepdown F
128 (REGW-F) test ($p \leq 0.05$), with the SPSS software Inc. (Chicago, United States).

129

130 **3. Results and discussion**

131 In order to prolong health and quality product, investigations on the effect of anti-
132 ethylene compounds on the disease development on cut flowers were performed. All the
133 1-MCP treatments significantly slowed down the development of grey mould, compared
134 to the inoculated control. The not inoculated control did not show any disease symptom
135 during the experiment, indicating the health of the plant material used. Significant
136 differences ($p \leq 0.05$) were denoted among the three 1-MCP applications/concentrations
137 (Fig. 1). Treatment with the lower CD-NS complex concentration (0.25 $\mu\text{L L}^{-1}$)
138 performed similarly or better than the commercial gaseous 1-MCP until day 13. At day
139 14, the lower dose of CD-NS complex resulted the best effective. At day 15, the
140 pathogen infection reached 100% value in all the inoculated flowers. Application of the
141 higher NS complex concentration (0.5 $\mu\text{L L}^{-1}$) was less active than the other two 1-MCP
142 treatments. Seglie et al. (2011a) already reported a lower activity of the higher
143 concentration of NS complex in extending the vase life of carnation cut flower. This
144 result might be explained by considering that the increase of the total amount of
145 nanosponge decreases the antagonist release (Seglie et al., 2011b).

146 Data about endogenous ethylene production were strictly related to the development
147 of grey mould on flowers. The lowest ethylene production was measured in flowers
148 treated with 0.25 $\mu\text{L L}^{-1}$ NS complex (0.53 $\mu\text{L L}^{-1}$), followed by gaseous 1-MCP-treated

149 flowers ($0.70 \mu\text{L L}^{-1}$). The not inoculated control and the flowers treated with the higher
150 concentration of NS complex produced the same ethylene concentration ($1.15 \mu\text{L L}^{-1}$).
151 As expected, the highest endogenous ethylene production was observed in the
152 inoculated control ($1.77 \mu\text{L L}^{-1}$) (Fig. 2). It could be assumed that NSs are able to
153 reduce ethylene production, by slowly releasing the ethylene antagonist, reducing the
154 senescence process, and, simultaneously, by adsorbing the phytohormone, and other
155 trap targeted organic compounds (Li and Ma, 1999). Ethylene antagonists could
156 maintain membrane integrity of plant tissues (Elad, 1997), by reducing *Botrytis* blight
157 of rose (Elad, 1995) and other plants (Elad et al., 1993). Anyway, the relationship
158 between ethylene synthesis, pathogen infection, and ethylene antagonists, such as NS
159 complex, should be further elucidated.

160 In conclusion, 1-MCP included in nanosponges can be a promising formulation to be
161 developed to control fungal diseases of cut flowers in the postharvest environment,
162 though the mechanism of action needs further elucidation.

163

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207

208 **Figure captions**

209 **Fig. 1** Effect of two different concentrations (0.25 and 0.5 $\mu\text{L L}^{-1}$) of β -CD-based-
210 nanosponge 1:8 - 1-MCP complex (NS complex) on grey mould development of
211 carnation cut flowers, compared to commercial gaseous 1-MCP (0.25 $\mu\text{L L}^{-1}$ for 6 h),
212 and to an inoculated (B^+) and not inoculated (B^-) controls. Vertical bars show the
213 confidence intervals (95%) of mean values.

214 *Mean separation within columns by the Ryan-Einot-Gabriel-Welsch's multiple
215 stepdown F (REGW-F) test, $p \leq 0.001$.

216

217 **Fig. 2** Endogenous ethylene production in cut flowers of *Dianthus caryophyllus* 'Idra di
218 Muraglia'. Flowers were treated with β -CD-based nanosponge 1:8 - 1-MCP complex
219 (NS complex; 0.25 and 0.5 $\mu\text{L L}^{-1}$, a. i.), and gaseous 1-MCP treatment (0.25 $\mu\text{L L}^{-1}$ for
220 6 h). Controls were inoculated (B^+) or not (B^-) with *Botrytis cinerea*.