



Qualitative responses of rocket varieties to biostimulants application

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Introduction

Nutritional quality and health benefit of vegetables are often related to the accumulation of biological active substances like secondary metabolites and phytonutrients. Numerous scientific evidences demonstrated the positive impact of plant biostimulants on different product quality traits (Bulgari et al., 2019). Rocket salad includes two important genera, *Diplotaxis* and *Eruca*, particularly appreciated for their peculiar flavour and the relevant content of diverse phytochemicals (Bell and Wagstaff, 2019). This work aims to investigate the effect of two commercial biostimulant products on some quality traits in two cultivars of *Eruca sativa* Mill. and *Diplotaxis tenuifolia* (L.) DC.

Materials & Methods

Location & Season	Cultivation system	Determinations	Species	Biostimulant products
Cornaredo, Milan (Italy) Summer 2021 (June – July)	Field (plastic tunnel greenhouse)	NFI, Flavonol (<i>in vivo</i> MPM-100) Sucrose, Nitrate (Spectrophotometric)	<i>Diplotaxis tenuifolia</i> (L.) DC (D1 –D2) <i>Eruca sativa</i> (Mill.) (E1-E2)	Megafofol® 0.3 mL/L Actiwave® 3 mL/L

Results

Flavonol content was higher in *Eruca* cultivars than in *Diplotaxis* cultivars (Tab. 1); in particular, the values ranged between 0.80 and 1.03 (F660nm / F325nm) and between 0.50 and 0.82 (F660nm / F325nm), respectively. Moreover, different trends were observed in response to the biostimulants applications. D2 plants treated with Actiwave® showed a higher content of flavonol than D2 control, whereas Megafofol® treatment did not induced any changes in the same cultivar. On the contrary, D1 plants treated with Megafofol® had a high level of flavonol if compared with D1 plants treated with Actiwave®.

Tab. 1 Flavonol content measured *in vivo* in rocket leaves

Species/Cultivar	Flavonol content (F660nm / F325nm) ^a		
	Control	Actiwave®	Megafofol®
<i>Diplotaxis tenuifolia</i>			
D1	0.64 ± 0.045abB	0.50 ± 0.013bC	0.67 ± 0.040aB
D2	0.58 ± 0.053bB	0.82 ± 0.049aB	0.71 ± 0.038abB
<i>Eruca sativa</i>			
E1	0.87 ± 0.052aA	0.88 ± 0.029aAB	0.80 ± 0.043aB
E2	0.98 ± 0.027aA	1.03 ± 0.038aA	0.97 ± 0.033aA

^a Values are means ± SE (n=6). Different lowercase letters indicate significant differences within cultivars, different uppercase letters indicate significant differences within treatments, according to the Tukey post-hoc test (p ≤ 0.05).



Fig. 1 Pictures of rocket during cultivation

The levels of sucrose ranged between 150.6 and 248.9 mg kg⁻¹ FW in *Diplotaxis* cultivars and between 107.7 and 232.1 mg kg⁻¹ FW in *Eruca* cultivars (Tab. 2). All rocket cultivars had similar concentration of sucrose, regardless the biostimulant application. A significant decrease was observed only in E1 plants treated with Megafofol®. This might be due to a translocation or a rapid metabolism of the sugar inside plants after the treatment. Figure 2 shows the NFI of rocket species/cultivars calculated from flavonols and chlorophyll content measured *in vivo* at harvest. The NFI was higher in *Diplotaxis* cultivars than in *Eruca* cultivars; Moreover, the four cultivars showed different trends in response to biostimulants applications.

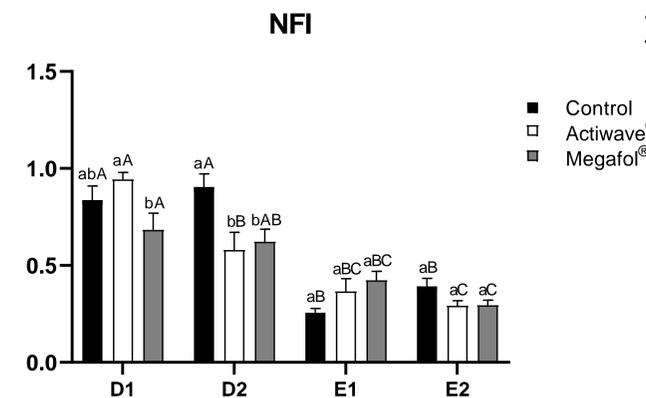


Fig. 2 Nitrogen flavonol index (NFI)

Tab. 2 Sucrose concentration in rocket leaves

Species/Cultivar	Sucrose (mg kg ⁻¹) ^a		
	Control	Actiwave®	Megafofol®
<i>Diplotaxis tenuifolia</i>			
D1	198.7 ± 29aA	150.6 ± 11aA	158.5 ± 17aA
D2	175.2 ± 5aA	248.9 ± 31aA	184.1 ± 21aA
<i>Eruca sativa</i>			
E1	221.7 ± 33aA	171.1 ± 4abA	107.7 ± 32bA
E2	232.1 ± 32aA	158.9 ± 29aA	167.9 ± 32aA

^a Values are means ± SE (n=3). Different lowercase letters indicate significant differences within cultivars, different uppercase letters indicate significant differences within treatments, according to the Tukey post-hoc test (p ≤ 0.05).

Conclusions and perspectives

Results revealed a variability in the quality traits depending on the combined effect of biostimulants application and genetic background. The present data will be integrated with additional analyses in order to better understand the responses observed in *Diplotaxis* and *Eruca* cultivars after the biostimulants applications. Further investigations will help to clarify the results obtained in the present experiment and to find which biostimulant product is better for the specific varieties.