

INTRODUCTION

- Ethyl formate (EF) is an FDA-approved food-flavoring agent with a GRAS status.
- It is a potent insecticide and antimicrobial promising as an alternative to the fumigant, methyl bromide.
- However, EF is highly volatile, flammable, and susceptibility to hydrolytic degradation, make its end-use application very challenging.

OBJECTIVES

- Development of EF precursor with enhanced storage stability.
- Encapsulation of EF precursor in electrospun fibers.
- Controlled release of EF for active packaging application.

METHOD

- Precursor was synthesized, which can be hydrolyzed to activate the release of EF vapor.

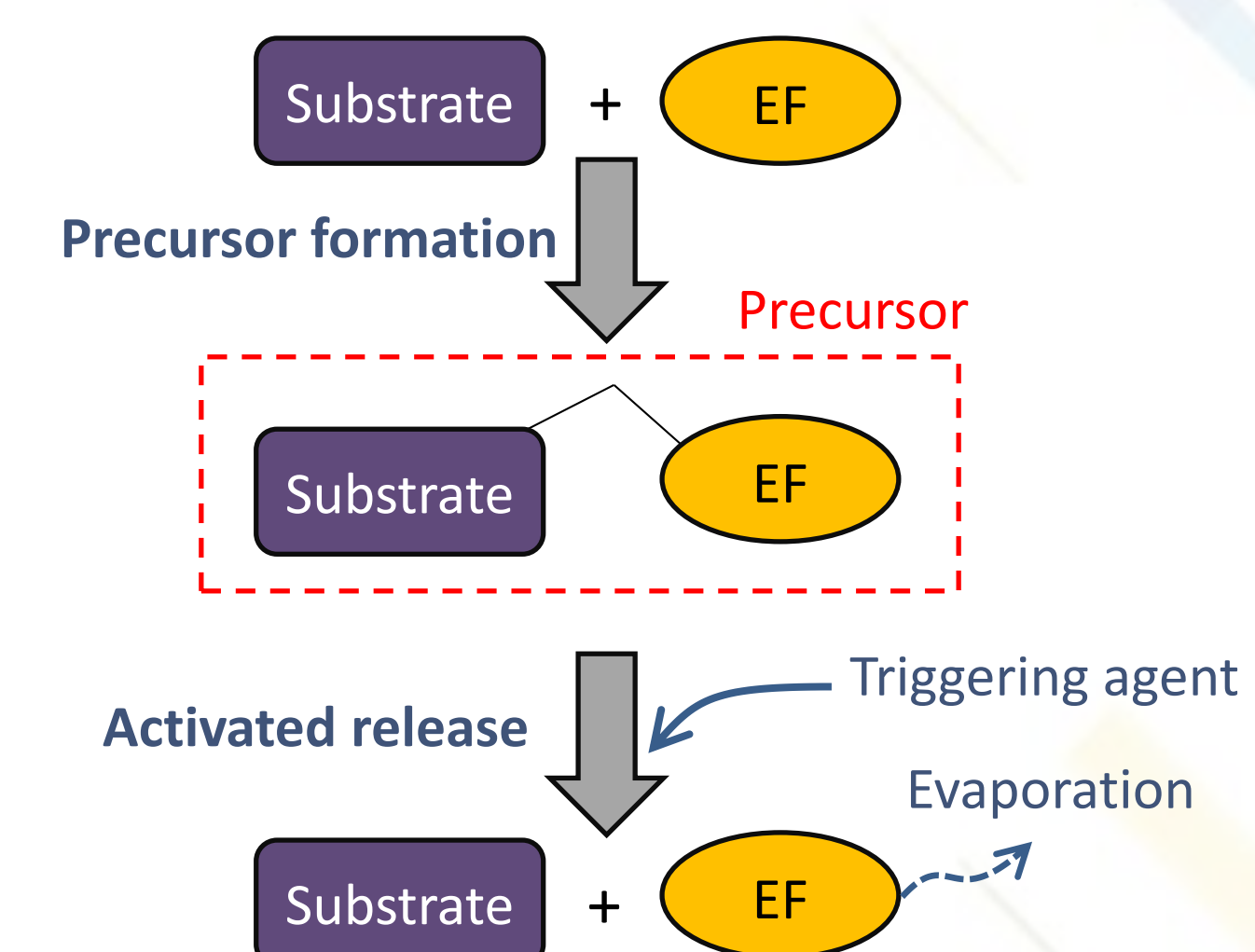


Fig. 1: Formation of precursor of EF and its release through cleavage of covalent bonds by a triggering agent.

- Scanning electron microscopy was used to study the morphologies of the fibers, which were correlated with the rheological properties of the spin dope solutions.
- EF release was triggered using 0.1 N citric acid. The release kinetic of EF from the fiber was studied using gas chromatography at 5, 10, 15, and 25°C (Fig. 3).

- The precursor was encapsulated in ethyl cellulose/poly(ethylene oxide) (EC-PEO) nonwoven by electrospinning (Fig 2).

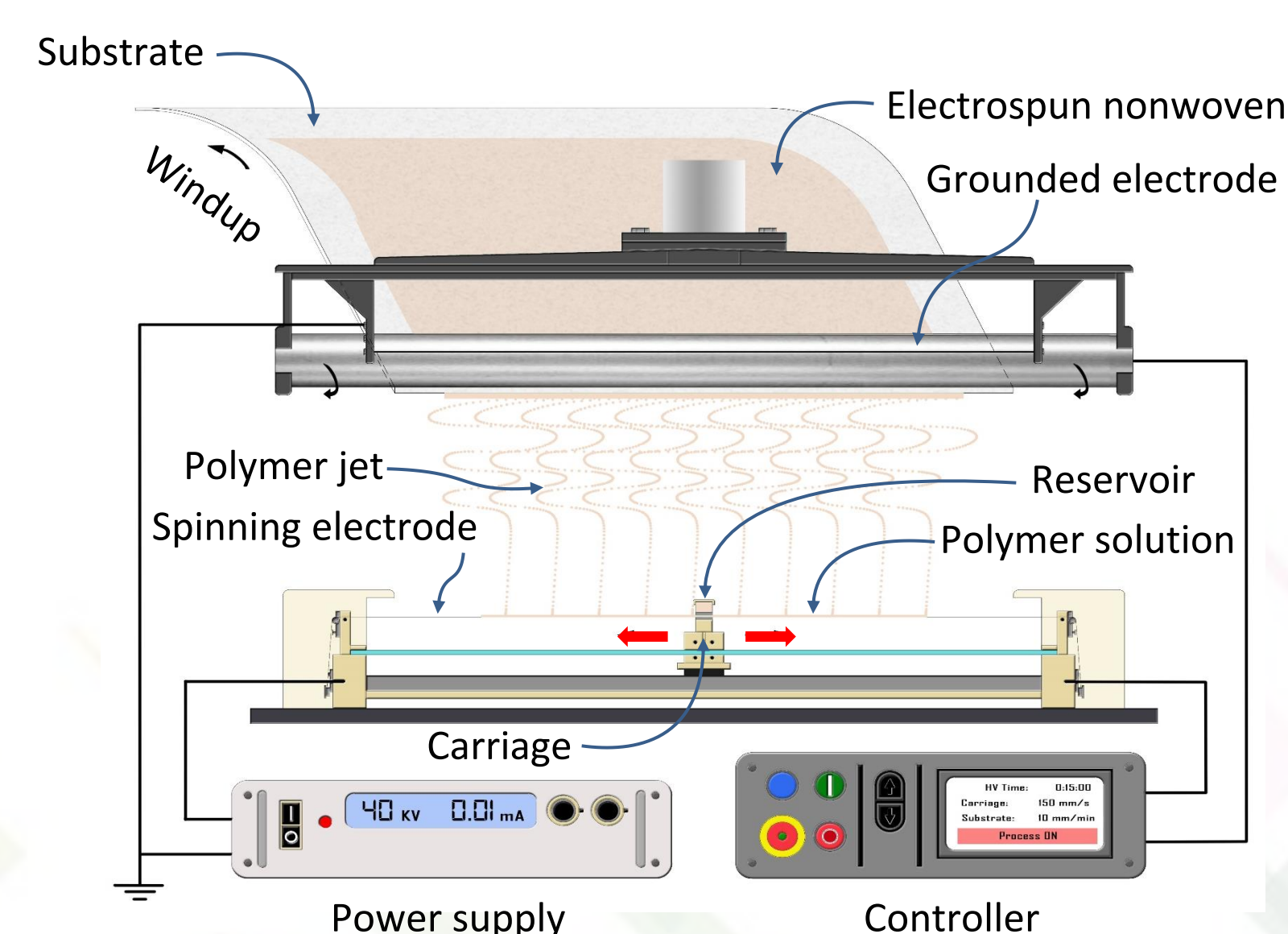


Fig. 2: Schematic of free surface wire electrospinner

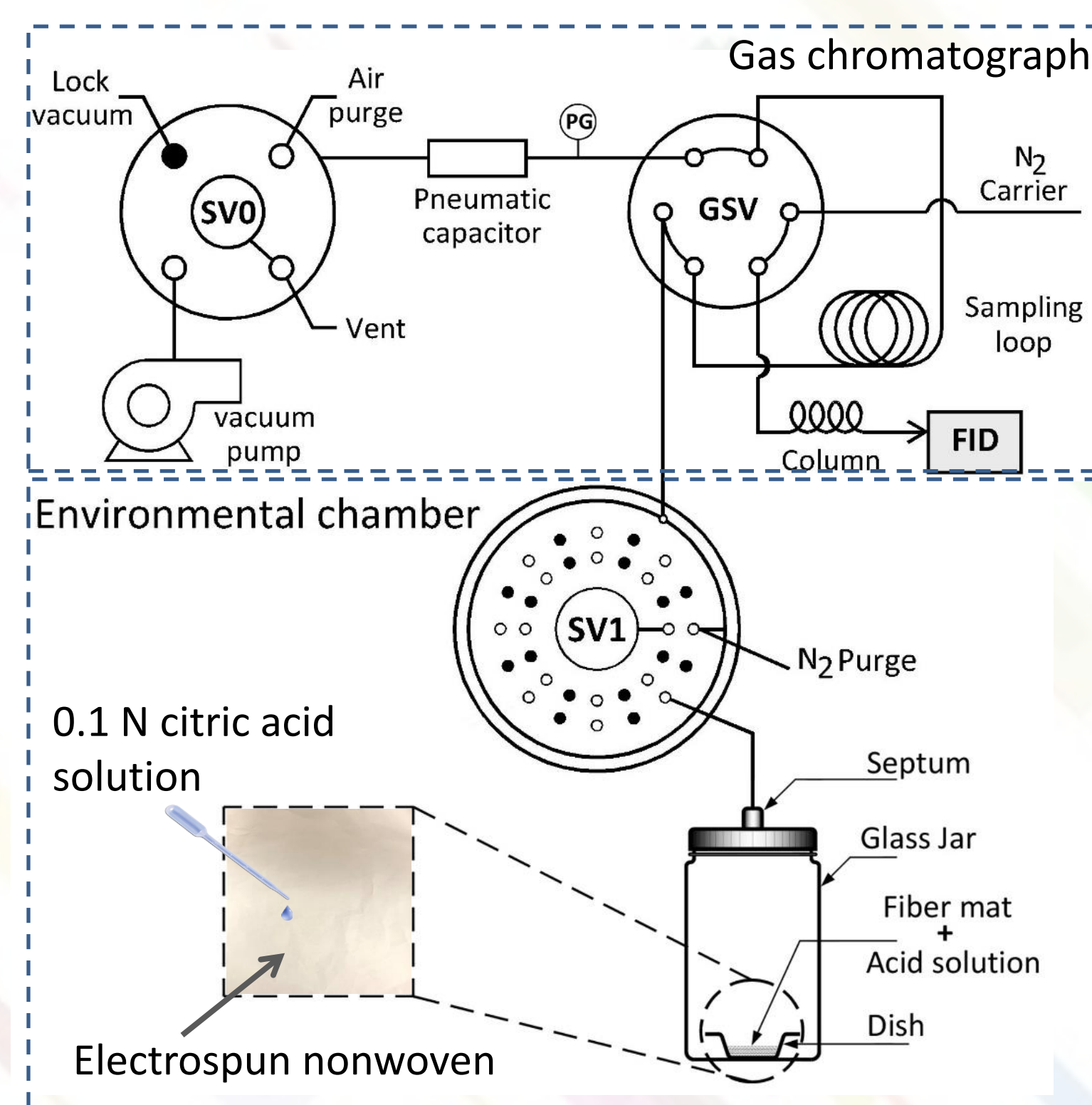


Fig. 3: Schematic representation of the automatic headspace analysis system used for studying the EF release.

RESULTS

- All spin dope solutions exhibited shear-thinning behaviors.
- The addition of EF precursor caused an increase in the solution apparent viscosity (Fig. 4), which correlated with the larger fiber diameter for precursor-loaded nonwovens than the pristine counterpart (Fig. 5).
- Increasing the temperature significantly increased the release of EF from the precursor-loaded fibers (Fig. 6).

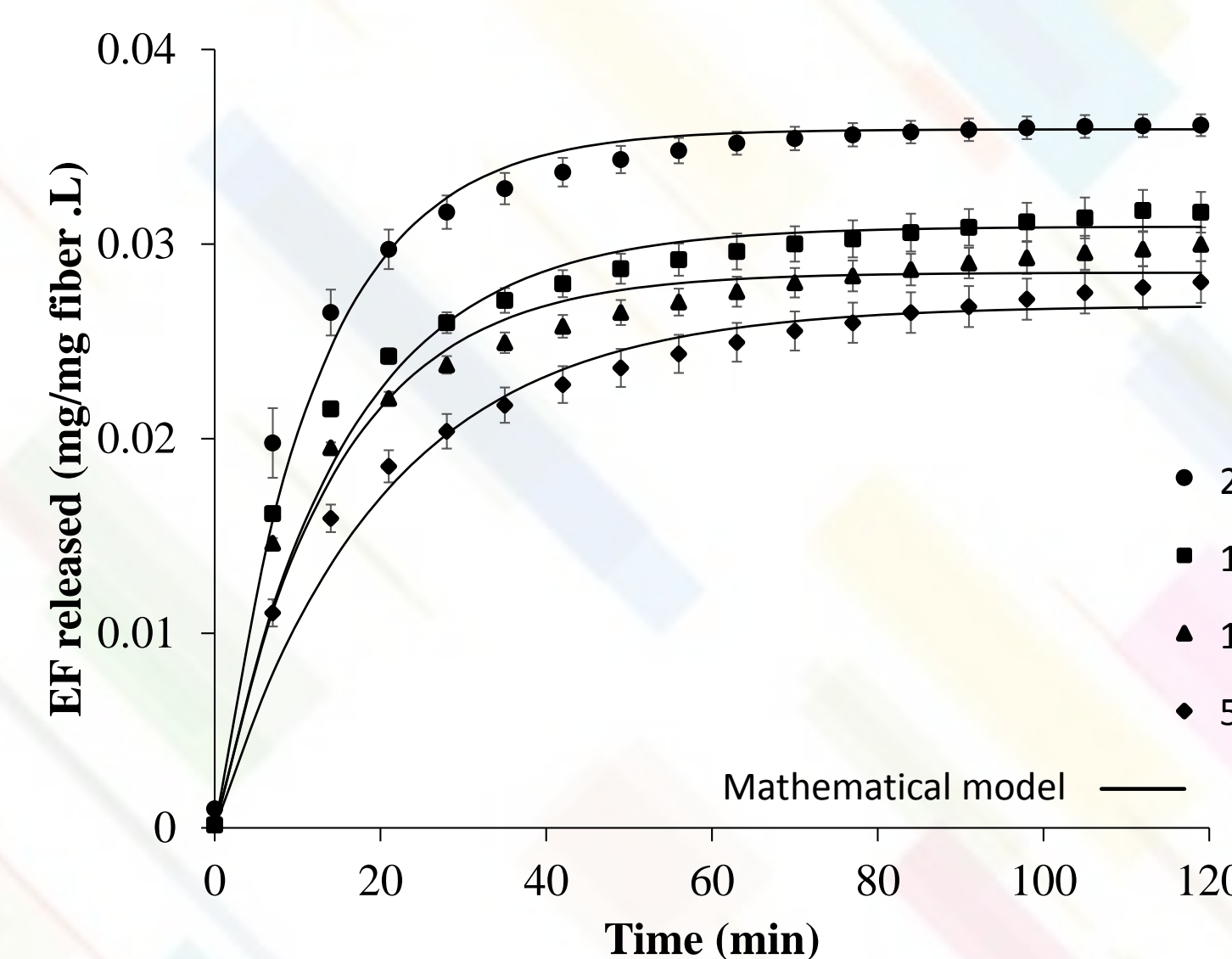


Fig. 6: EF released from the precursor-loaded fibers at different temperatures using 0.1 N citric acid solution.

Table 1: EF released from the fiber at different temperatures, and the fitted model parameters.

Temperature °C	EF released after 2 h		C _e mg/mg fiber. L	K min ⁻¹	R ²
	mg/mg fiber. L	%			
5	0.028	74	0.027	0.05	0.97
10	0.029	79	0.029	0.07	0.97
15	0.31	83	0.031	0.075	0.97
25	0.036	95	0.036	0.082	0.99

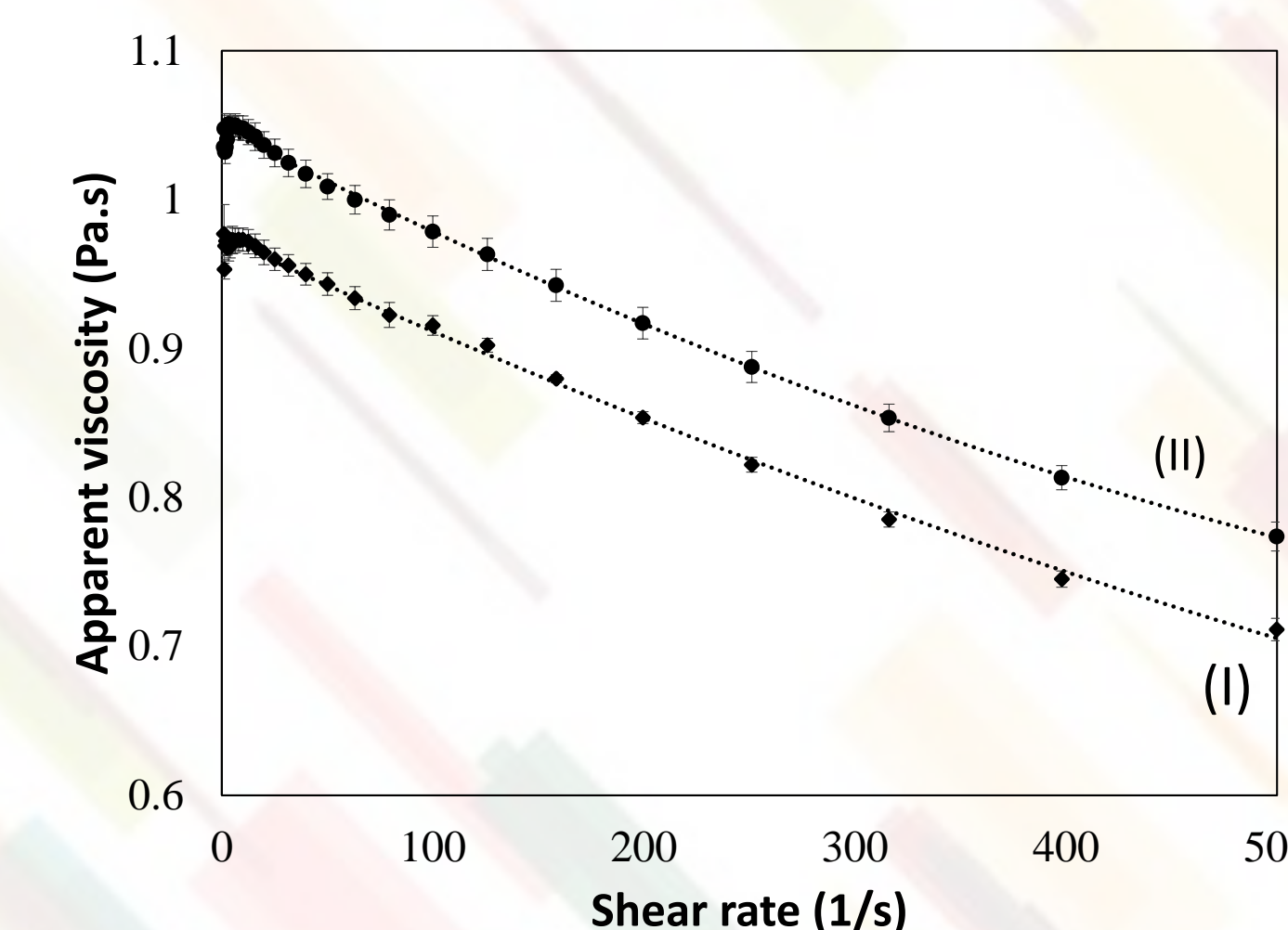


Fig. 4: Apparent viscosity of EC-PEO-doped solution (I) and precursor-doped solution (II).

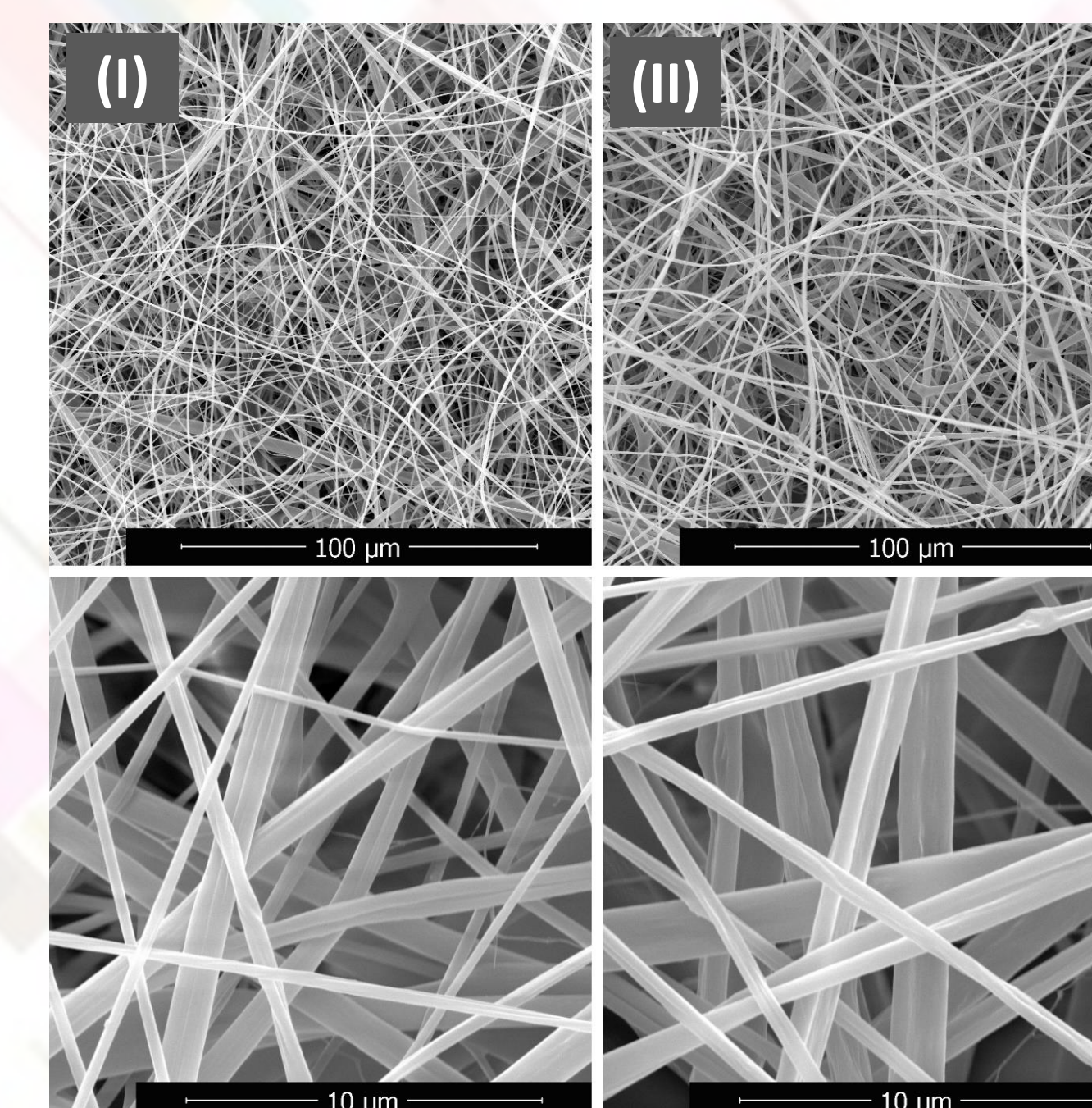


Fig. 5: SEM micrographs and histograms of fiber diameter distributions of pristine fiber (I) and precursor-loaded fiber (II).

CONCLUSIONS

- EF precursor was synthesized for activated release of EF vapor.
- Electrospun EC-PEO nonwoven was developed as a carrier for the EF precursor.
- 95 % of EF was released from the nonwoven after 2 h at 25°C.
- The approach of converting the highly volatile EF into a solid-state precursor can be useful for in-package fumigation of fresh produce to destroy insect pests, as well as to inhibit the proliferation of spoilage microorganisms.

ACKNOWLEDGEMENTS

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