

Low Temperature Polyethylene Biodeterioration by Freshwater Microplastics-Associated Bacteria

Abdoullah Hleihel^{1,2*}, Jonas M. Stadfeld^{2,*}, Kira L. Goff¹, Amelia M. Danzinger², Craig Beaver³, Troy Stuart³, Shauna Reckseidler-Zenteno¹, Janelle M. Baker⁴, Srijak Bhatnagar^{1,2}

Affiliations:

¹Faculty of Science and Technology, Athabasca University, Athabasca, AB, Canada

²Department of Biological Sciences, University of Calgary, Calgary, AB, Canada

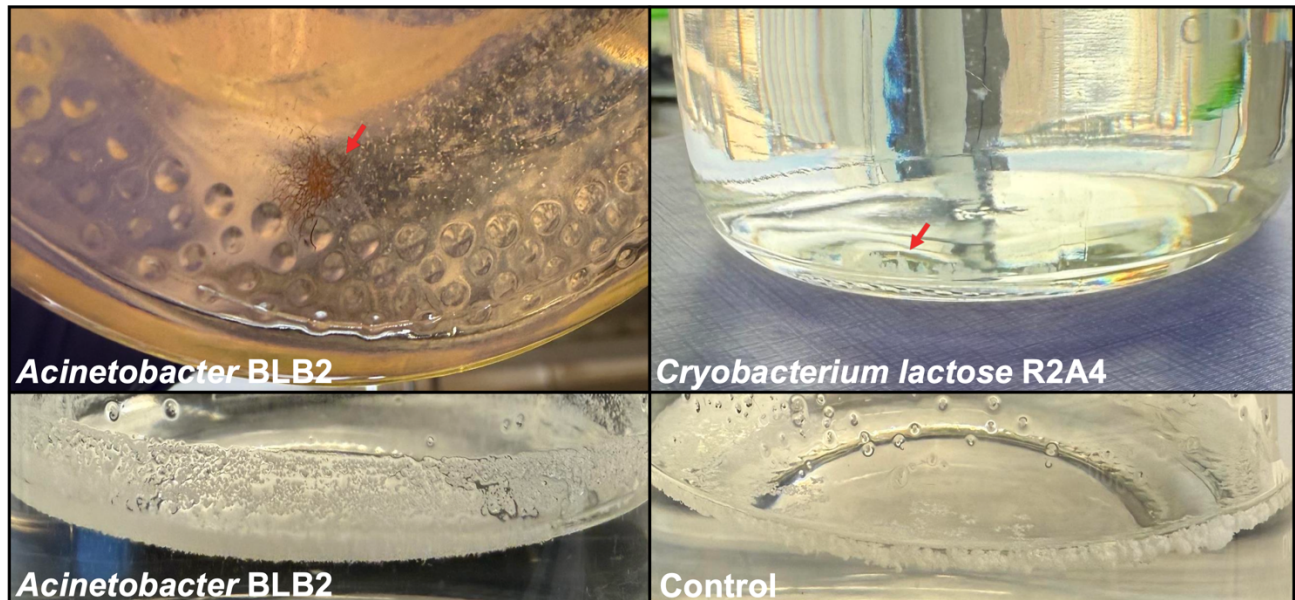
³Lands and Resources Department, Bigstone Cree Nation, Wabasca, AB, Canada

⁴ Faculty of Humanities and Social Sciences, Athabasca University, Athabasca, AB, Canada

*Authors contributed equally

Corresponding Author: Srijak Bhatnagar, sbhatnagar@athabascau.ca

16 **Supplementary Information**



Supplementary Figure 1. Visual observations of LDPE degradation microcosms.

(a) *Acinetobacter* BLB2 showing green-brown web-like growth (b) *Cryobacterium* BR2A4 exhibiting translucent floating aggregates (c) *Acinetobacter* BLB2 microcosm highlighting visibly smaller LDPE fragments with increased bulk dispersion compared to (c) control microcosm

18
19 **Supplementary Data 1.** Metadata of the 14 Isolates used for the LDPE degradation
20 experiment.

21 **Supplementary Data 2.** Final calculated weight losses of the 14 isolates and
22 experimental control after a seven month incubation period.

23 **Supplementary Data 3.** Monthly optical Density (600nm) measurements of microcosms
24 during LDPE degradation experiment.

25 **Supplementary Data 4.** Cell densities (cells/mL) of the 14 isolates including standard
26 deviation at months five and six of incubation.

27 **Supplementary Data 5.** GCMS results of detected compounds, their assigned
28 classification, and concentration ($\mu\text{g/L}$) of each based on a C16 response factor after a
29 seven month incubation period.