1	SUPPLEMENTARY MATERIAL for				
2	Polyvinyl chloride degradation by a bacterium isolated from the				
3	gut of insect larvae				
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# 29 Supplementary Tables and Figures

# 30 Supplementary Table 1 Detection and analysis results of additives in PVC film by GC-

31 MS analysis

Name	RT/min	Qualitative and quantitative transition	Standard curve/R <sup>2</sup>	Content in PVC film (%)
DOA	11.78	129,57,112/129	y=2E+06x-226835, R <sup>2</sup> =0.9925	22.90
DOTP	14.53	70,112,149/70	y=952890x-2E+06, R <sup>2</sup> =0.9935	5.23
Erucylamide	15.09	59,72,55/59	y=523400x-1E+06, R <sup>2</sup> =0.9936	2.05

# Supplementary Figure 1 Characterization of morphological changes of PVC film in the feces of *Spodoptera frugiperda* larva

a, Scanning electron microscopy (SEM) results showing clear degradation of PVC film
recovered from larval feces in PVC group. b, SEM results showing no visible biodegradation
of PVC film recovered from feces in Antibiotic group (in which gentamicin antibiotic was
used to inactivate most gut microbes of the larva). For both a and b, subfigure (1) to (4)
represented the degradation of PVC film from low magnification to high magnification in SEM.
At least 3 times experiment was repeated independently.



# 42 Supplementary Figure 2 Screening and isolation of PVC film degrading strains and 43 phylogenetic analysis of 16S rRNA gene sequence

a, Experimental screening and isolation (The dissected intestine of larva in PVC group was
used to inoculate and enrich for degrading strain EMBL-1, which was cultured on PVC film
in MSM liquid medium and LB solid medium). b, Phylogenetic tree of 16S rRNA gene
sequences showing PVC-degrading strain EMBL-1 as a novel Klebsiella strain most closely
related to *Klebsiella variicola* and *Klebsiella pneumoniae* (The analysis was conducted using
the Fast Minimum Evolution method at a maximum sequence difference of 0.75).

a





## 51 Supplementary Figure 3 Degradation results of strain EMBL-1 on PVC film

- a, SEM images of the cracks and pits formed on the PVC film by strain EMBL-1 on day 90.
- 53 At least 3 times experiment was repeated independently. b, Water contact angle of PVC film
- 54 after co-culturing with strain EMBL-1 significantly decreased compared with the control
- group (t-test P < 0.05, n = 3 samples each). c, Tensile strength of PVC film in the control group
- and the EMBL-1 group after 90 days. d, OD<sub>600</sub> values during 65-day growth of strain EMBL-
- 1 using soybean oil (SO) as the sole organic carbon substrate, which showed no growth of
- strain EMBL-1 compared with the control groups. Each group had 3 replicates and the mean
- <sup>59</sup> values were visualized. e-f, Thermal gravimetric analysis (TGA) results of PVC film in control
- and EMBL-1 groups during 90 days. g, FTIR results of PVC film in control groups during 90
- 61 days. Source data are provided as a Source Data file.



## 63 Supplementary Figure 4 Spectrum of NMR experiments of PVC

a, DOSY Spectrum of pure PVC in control and EMBL-1 group on 90 days. b, 2D <sup>1</sup>H-<sup>1</sup>H COSY
spectrum of pure PVC in EMBL-1 group on 90 days. c, 2D <sup>1</sup>H-<sup>1</sup>H NOESY spectrum of pure
PVC in EMBL-1 group on 90 days. d, 2D <sup>1</sup>H-<sup>13</sup>C HSQC spectrum of pure PVC in EMBL-1
group on 90 days. e, 2D <sup>1</sup>H-<sup>13</sup>C HMBC spectrum of pure PVC in EMBL-1 group on 90 days.



- 71 Supplementary Figure 5 Detection results of degradation products of PVC film by GC-
- 72 **MS**
- a, the TIC diagram of degradation products of PVC film in two groups by the time of 90 d. b-
- f, the mass spectrum and structural formula of the potential degradation products 1-5.



## 77 Supplementary Figure 6 Detection and identification results of additives in PVC film and

## 78 the degradation activity of strain EMBL-1 on main additives

a, the Py/GC-MS diagram of three additives in PVC film. b, the TIC diagram of three additives
identification in PVC film. c-e, the mass spectrum of three additives. f, the result of
degradation activity of strain EMBL-1 on three additives. The experiments were conducted in
triplicates, and mean value ± standard deviation was visualized.



#### 84 Supplementary Figure 7 The whole-genome map of strain EMBL-1



### Metabolism

- [C] Engergy production & conversion
- [G] Carbohydrate transport & metabolism
- E] Amino acid transport & metabolism
- F] Nucleotide transport & metabolism
- [H] Coenzyme transport & metabolism
- [I] Lipid transport & metabolism
- [P] Inorganic ion transport & metabolism
- [Q] Secondary metabolites biosynthesis, transport & catabolism

### Information storage & processing

- [J] Translation, ribosomal structure & modification
- [A] RNA Processing & modification
- [K] Transcription
- [L] Replication, recombination & repair
- [B] Chromatin structure & dynamics

### Cell processing & signaling

- [D] Cell cycle control , cell division, chromosome partitioning
- [Y] Nuclear structure [V] Defense mechanisms
- [T] Signal transduction mechanism
- [M] Cell wall/membrance/envelope biogenesis
- [N] Cell motility
- [Z] Cytoskeleton
- [W] Extracellular structures
- [V] Intracellular trafficking, secretion & vesicular transport
   [O] Posttranslational modification, protein turnover, chaperones

## Poorly characterized

- [R] General function prediction only
- [S] Function unknown
- [X] No COG assignment

# 87 Supplementary Figure 8 Results of proteome experiments and expression purification of 88 catalase-peroxdiase

a, Growth curve of strain EMBL-1 under different carbon source conditions. b, The 89 concentration of proteins in four groups detected by Bradford method. c, Construction results 90 of expression strain of catalase-peroxdiase (M means protein marker,1 means the strain lysate 91 before induction, 2 means the strain lysate after induction ). d, purification results of catalase-92 peroxdiase using Ni-IDA Agarose Magnetic Beads (M means protein marker,1 means the 93 strain lysate after induction, 2-3 mean the eluention during purification, 4 means the eluention 94 with catalase-peroxdiase). For a and b, the experiments were conducted in triplicates, and mean 95 value  $\pm$  standard deviation was visualized. For c and d, the experiments were repeated at least 96 97 3 times independently. Source data were provided as a Source Data file.

98



# Supplementary Figure 9 Physicochemical characterization of PVC degradation by catalase-peroxidase

a, FTIR results for PVC in the PVC-Cp group and PVC control group. b, Molecular weight of
PVC in the PVC-catalase-peroxidase group and PVC control group. The experiments were
conducted in triplicates and mean value ± standard deviation was visualized. Source data were
provided as a Source Data file.

107



110 Supplementary Figure 10 Potential degradation products detection of PVC by catalase-

# 111 peroxidase

- a, The detection results of GC-MS of degradation products from PVC-catalase-peroxidase
- 113 group and PVC control group. b, the mass spectrum and structural formula of the potential
- 114 degradation products.

