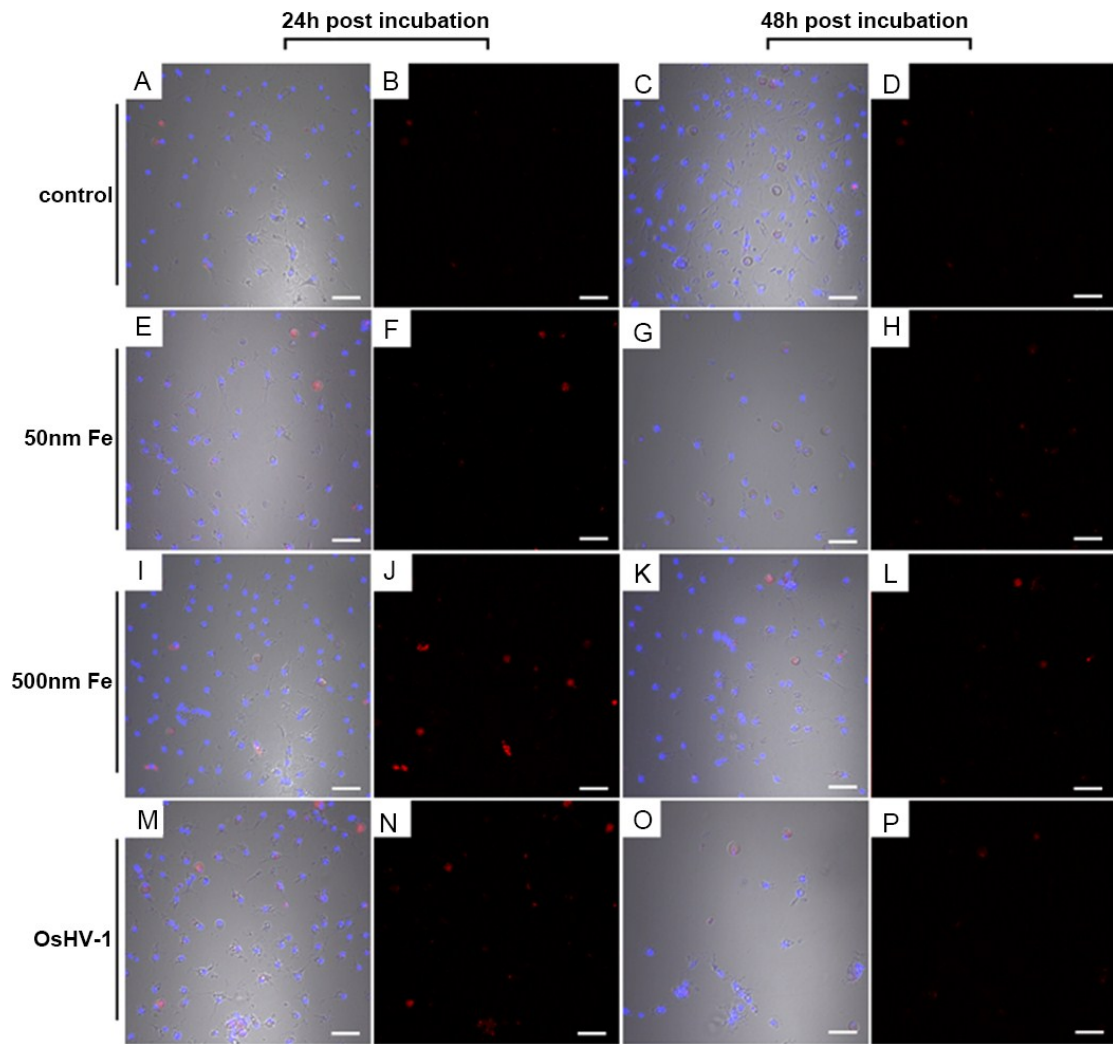


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2 Fig.S1. Transmission electron microscopy confirmation of OsHV-1 infection in ark
3 clams. Tissues, (A) Hemocytes, (B) Mantle, (C) Hepatopancreas, were examined at 72
4 hpi. Arrows indicate OsHV-1 particles.

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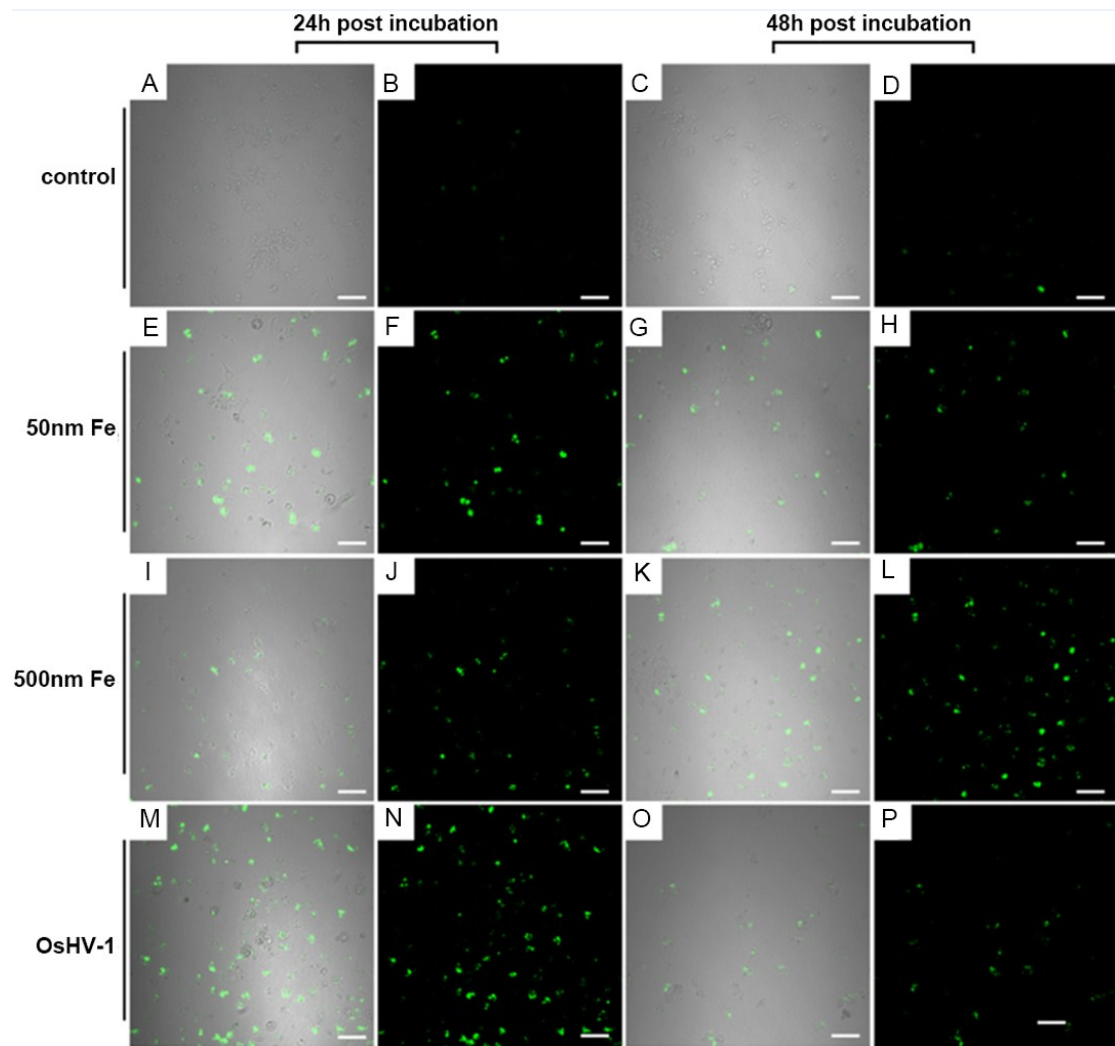
7

8 Fig.S2. ROS generation of in vitro hemocytes was promoted under incubation with high
 9 level iron and OsHV-1. (A-D) Normal cultured hemocytes were set as the control
 10 group. Few ROS signal (red) could be detected. (E-H), (I-L) Hemocytes under
 11 incubation with supplemented 50 nm and 500 nm iron respectively. More ROS signal
 12 (red) was found. The amount of adherent hemocytes decreased after 72 h incubation
 13 with supplemented 50 nm and 500 nm iron. Hemocytes incubated with OsHV-1 also
 14 showed more ROS signal (red) (M-P). (scale bar = 50 μ m).

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20 Fig.S3. Hemocyte death in vitro was induced under incubation with high level iron and
 21 OsHV-1. (scale bar = 50 μm) (A-D) Normal cultured hemocytes were set as the control
 22 group. Few dying hemocyte signal (green) was detected. The increase of dying
 23 hemocyte signal (green) could be observed in (E-H), (I-L) Hemocytes under incubation
 24 with supplemented 50 nm and 500 nm iron respectively; and in (M-P) Hemocytes
 25 incubated with OsHV-1.

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28 **The promotion of ROS and cell death by the incubation with high level iron and**
29 **OsHV-1**

30 The fluorescent indicator 2',7'-dichlorofluorescein (DCF) was used to monitor the
31 levels of intracellular ROS in hemocytes under supplement iron incubation and OsHV-1
32 infection (Fig. S2). The results showed that the DCF fluorescence signals (red) were
33 enhanced at 24 h under iron incubation with 50 nM and 500 nM supplement iron, and
34 comparatively weakened from 24 h to 48 h with decreasing adherent hemocytes, which
35 was similar to the results under OsHV-1 infection. TUNEL assay were undertaken to
36 determine the effect of the incubation with supplementary iron and OsHV-1 on DNA
37 damage in hemocytes (Fig. S3). The results showed that the positive signals (green)
38 were promoted at 24 h under supplementary iron (50 nM and 500 nM) and OsHV-1
39 incubation. From 24 h to 48 h, the positive signal also weakened as the adherent
40 hemocytes decreasing.

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