Table S1: table displaying the effects of papaverine on oxidative stress as a change of fluorescence intensity relative to the fluorescence intensity of cells propagated in growth medium on MDA-MB-231 cells compared to A549- and DU145 cell lines at 48 h. Statistical significance is represented by an * when using the student t-test with a P value of 0.05 compared to cells propagated in growth medium

Cell line	Vehicle- treated cells	10 µM PPV- treated cells	50 µM PPV- treated cells	100 μM PPV- treated cells	150 μM PPV- treated cells	ESE-ol- treated cells
MDA-MB-231	1.00 ± 0.04	1.05 ± 0.07	0.91 ± 0.1	1.07 ± 0.08	1.11 ± 0.10	1.91 ± 0.04*
A549	0.95 ± 0.04	1.09 ± 0.08*	1.23 ± 0.06*	1.18 ± 0.06*	1.14 ± 0.04*	1.94 ± 0.04*
DU 145	1.00 ± 0.02*	0.92 ± 0.02*	0.90 ± 0.07	0.96 ± 0.04	0.99 ± 0.04	1.78 ± 0.06*

Cells propagated in growth medium

MDA-MB-231

ESE-ol-treated cells

Vehicle-treated cells

10 μ M PPV-treated cells

50 µM PPV-treated cells 100 µM PPV-treated cells 150 µM PPV-treated cells

Figure S1. Fluorescence staining showing H₂O₂ production in MDA-MB-231 cells after 48 h. Fluorescence microscopy images of DCFDA staining demonstrating the effects of PPV (10-150 µM) on the fluorescent intensity on MDA-MB-231 cells at 48 h at a magnification of x20. A scale bar of 200µM is included.



50 μM PPV-treated cells $\,$ 100 μM PPV-treated cells $\,$ 150 μM PPV-treated cells

Figure S2. Fluorescence staining showing H2O2 production in A549 cells after 48 h. Fluorescence microscopy images of DCFDA staining demonstrating the effects of PPV (10-150 µM) on the fluorescent intensity on A549 cells at 48 h at a magnification of x20. A scale bar of 200µM is included.

A549



50 μ M PPV-treated cells 100 μ M PPV-treated cells 150 μ M PPV-treated cells

Figure S3. Fluorescence staining showing H₂O₂ production in DU145 cells after 48 h. Fluorescence microscopy images of DCFDA staining demonstrating the effects of PPV (10-150 μ M) on the fluorescent intensity on DU145 cells at 48 h at a magnification of x20. A scale bar of 200 μ M is included.

Table S2: table displaying the effects of papaverine on oxidative stress as a change of fluorescence intensity relative to the fluorescence intensity of cells propagated in growth medium on MDA-MB-231 cells compared to A549- and DU145 cell lines at 72 h. Statistical significance is represented by an * when using the student *t*-test with a *P* value of 0.05 compared to cells propagated in growth medium.

Cell line	Vehicle- treated cells	10 µM PPV- treated cells	50 µM PPV- treated cells	100 μM PPV-treated cells	150 μM PPV-treated cells	ESE-ol- treated cells
MDA-MB- 231	0.97 ± 0.00*	1.04 ± 0.04	0.73 ± 0.02*	0.83 ± 0.03*	0.84 ± 0.01*	1.95 ± 0.02*
A549	0.98 ± 0.02	1.02 ± 0.03	0.92 ± 0.03*	0.75 ± 0.04*	0.69 ± 0.03*	1.94 ± 0.03*
DU 145	0.99 ± 0.01	1.44 ± 0.02*	1.05 ± 0.05	1.14 ± 0.04*	1.15 ± 0.02*	1.83 ± 0.03*



50 μM PPV-treated cells 100 μM PPV-treated cells 150 μM PPV-treated cells

Figure S4. Fluorescence staining showing H₂O₂ production in MDA-MB-231 cells after 72 h. Fluorescence microscopy images of DCFDA staining demonstrating the effects of PPV (10-150 μ M) on the fluorescent intensity on MDA-MB-231 cells at 72 h at a magnification of x20. A scale bar of 200 μ M is included.



50 μM PPV-treated cells 100 μM PPV-treated cells 150 μM PPV-treated cells

Figure S5. Fluorescence staining showing H₂O₂ production in A549 cells after 72 h. Fluorescence microscopy images of DCFDA staining demonstrating the effects of PPV (10-150 μ M) on the fluorescent intensity on A549 cells at 72 h at a magnification of x20. A scale bar of 200 μ M is included.



50 μ M PPV-treated cells 100 μ M PPV-treated cells 150 μ M PPV-treated cells

Figure S6. Fluorescence staining showing H₂O₂ production in DU145 cells after 72 h. Fluorescence microscopy images of DCFDA staining demonstrating the effects of PPV (10-150 μ M) on the fluorescent intensity on DU145 cells at 72 h at a magnification of x20. A scale bar of 200 μ M is included.