

SUPPLEMENTAL DATA

A.



B.

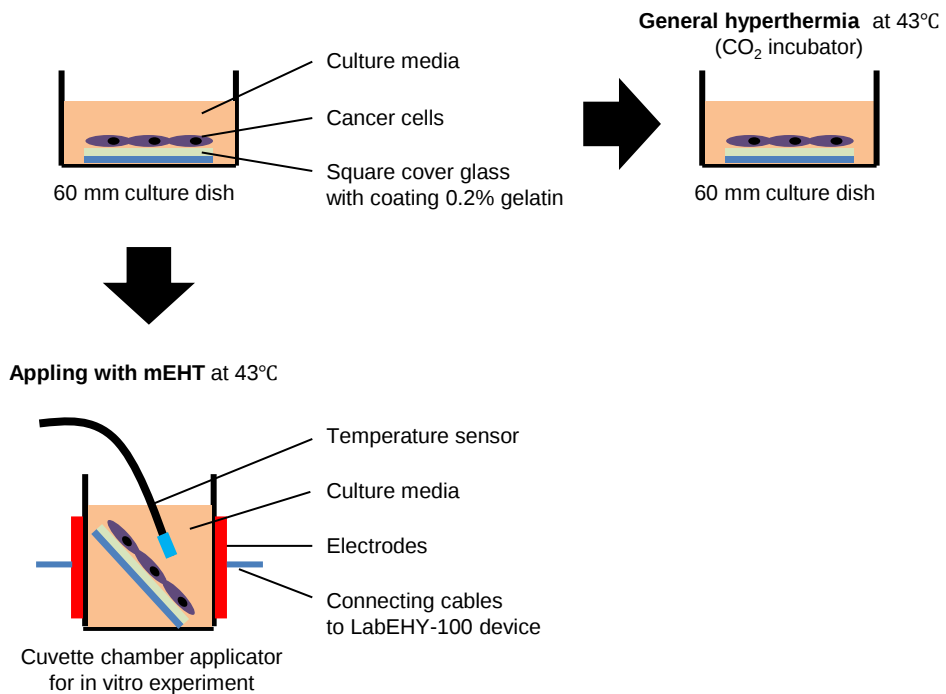
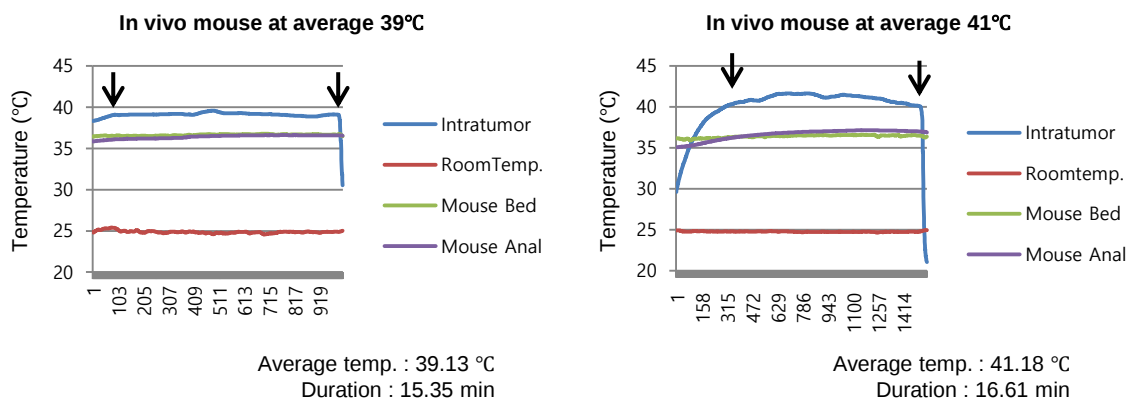


Figure S1. mEHT device for in vitro and in vivo experiment (LabEHY-100). (A) The LabEHY-100 is composed with an radiofrequency (RF) current modulator (①), 4 channel thermometer (②), connectors for 4 temperature sensors (③), in vivo mouse applicator with flexible textile electrode (④) and, connecting cables (⑤), including the suitable software which is monitored and recorded temperature and period (⑥). A suitable software collected the variation of radiofrequency, temperature change, and treated period and recorded these information. (B) Shown the diagram of in vitro cell experiment. Cancer cells were seeded in 0.2% gelatin coating cover glass and cultured at overnight (Top). Cover glass were placed at a cuvette chamber applicator and the energy of RF current were passed for a period of time (Bottom). General hyperthermia were increased temperature at 43°C using CO₂ incubator (Right).

A.



B.

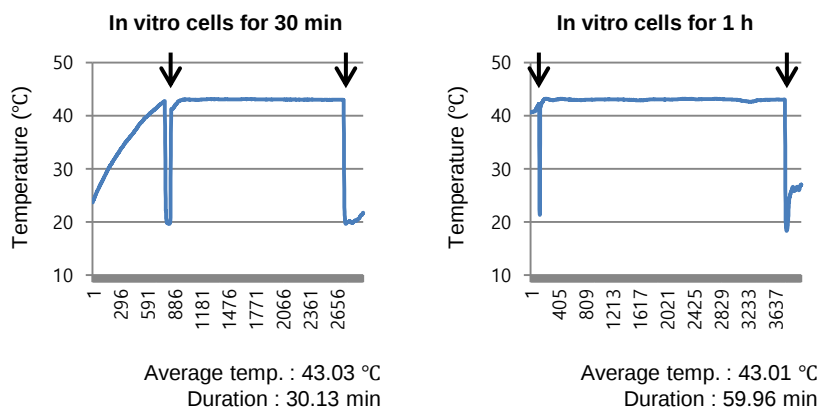


Figure S2. Representative graph of temperature change in mEHT treatment. Shown the representative graph of temperature and treated period. **(A)** xenograft tumors were applied with mEHT at 39°C (Left) or 41°C (Right) for 15 min. **(B)** cancer cells were applied at 43°C for 30min (Left) or 1 h (Right). Black arrow is start or end point of mEHT treatment.

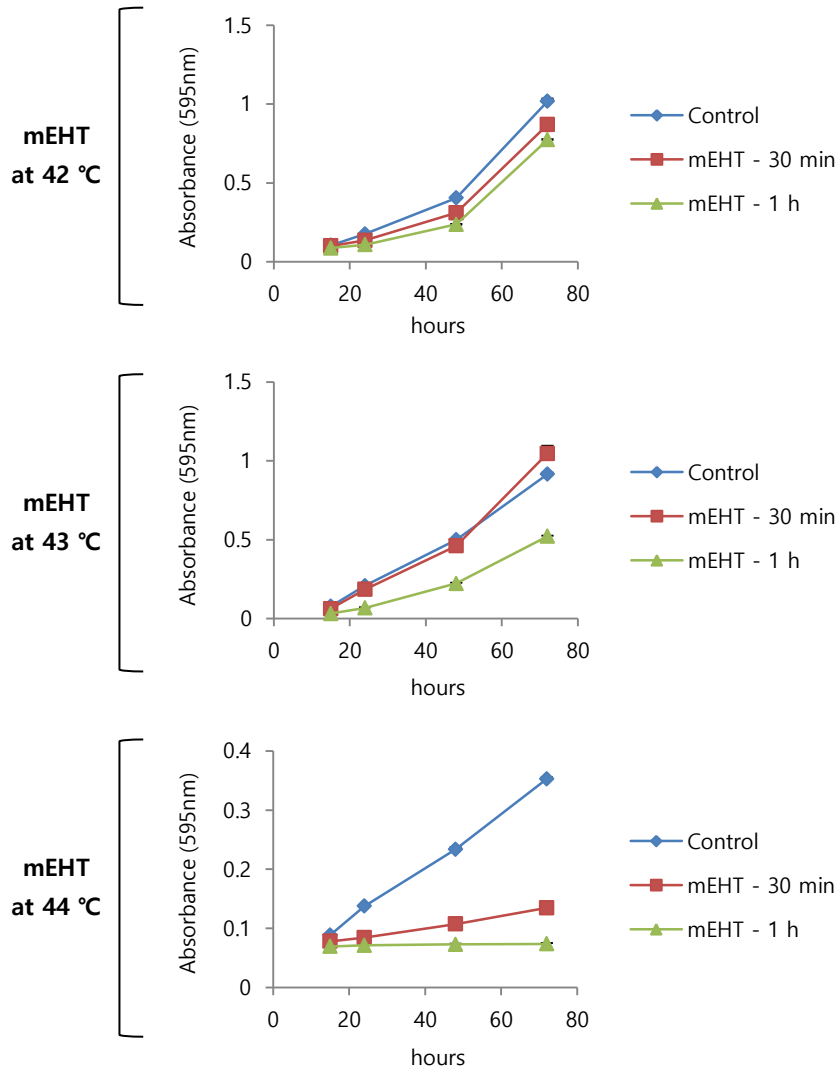


Figure S3. The effect of mEHT was evaluated in HeLa cells for optimizing experimental condition. 3×10^6 HeLa cells were seeded on 0.2% gelatin coated cover glass and were applied with mEHT at 42 °C, 43 °C and 44 °C for 30 min or 1 h. mEHT exposed HeLa cells were incubated for indicated time (16 h to 72 h) in the presence of 5% CO₂ at 37 °C in a incubator. Cell viability was measured via crystal violet assay. Results are the means $\pm$ S.E. n = 4.

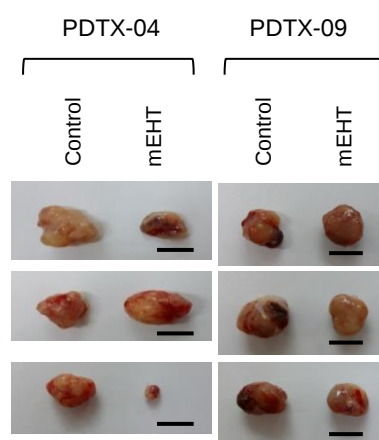
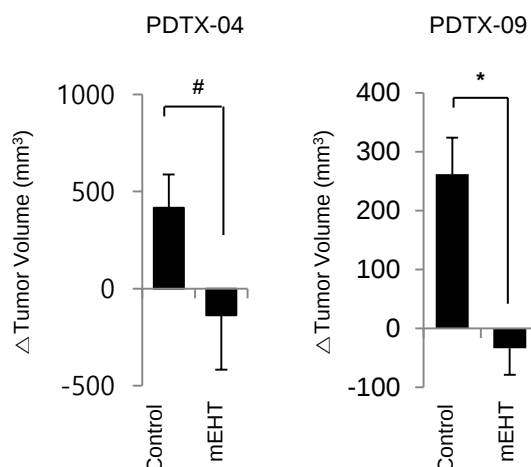
A.**B.**

Figure S4. mEHT suppressed the tumor growth in patients-derived tumor xenograft.

Athymic nude mice were transplanted subcutaneously into two patients-derived tumor xenograft (PDX) piece with roughly 3 mm in length and width. PDX-04 (Endometrial cancer) or PDX-09 (Ovarian cancer) tumors were applied with mEHT at 41°C for 15 min. (PDX-04, 5 times of mEHT between 35 and 44 day; PDX-09, 4 times of mEHT between 85 and 104 day). **(A)** Gross images of PDX tumors. Scale bar = 1 cm. **(B)** PDX tumor volume were measured using digital caliper. The box plot was shown the variation value of tumor growth. Results are the mean of growth variation (Δ tumor volume = tumor volume at end day – tumor volume at start day) $\pm$ S.E. $n = 3$. (#, $p > 0.05$; *, $p < 0.05$).

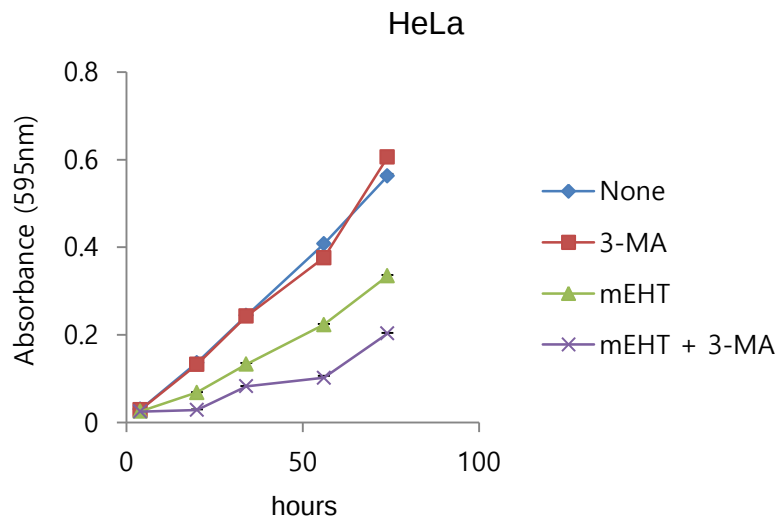
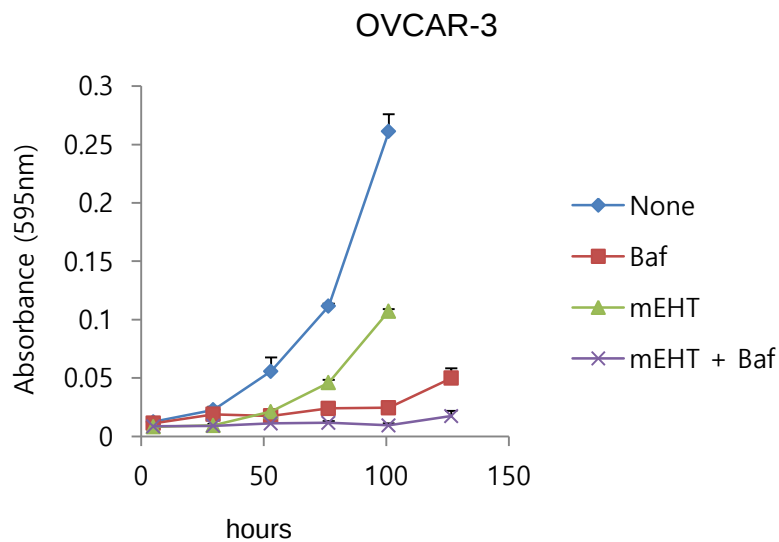
A.**B.**

Figure S5. Autophagy inhibitor enhanced apoptosis in mEHT-exposed HeLa and OVCAR-3 cancer cells. (A) HeLa cervical cancer cells were exposed to mEHT at 43°C for 1 h in the presence of 1 mM 3-methyladenine (3-MA). **(B)** OVCAR-3 ovarian cancer cells were exposed to mEHT at 43 °C for 1 h in the presence of 100 nM bafilomycin. mEHT exposed cells were incubated for 6 to 72 h at 37 °C and 5% CO₂ incubator. Cell viability was measured via crystal violet assay. Results are the means $\pm$ S.E. n = 4.

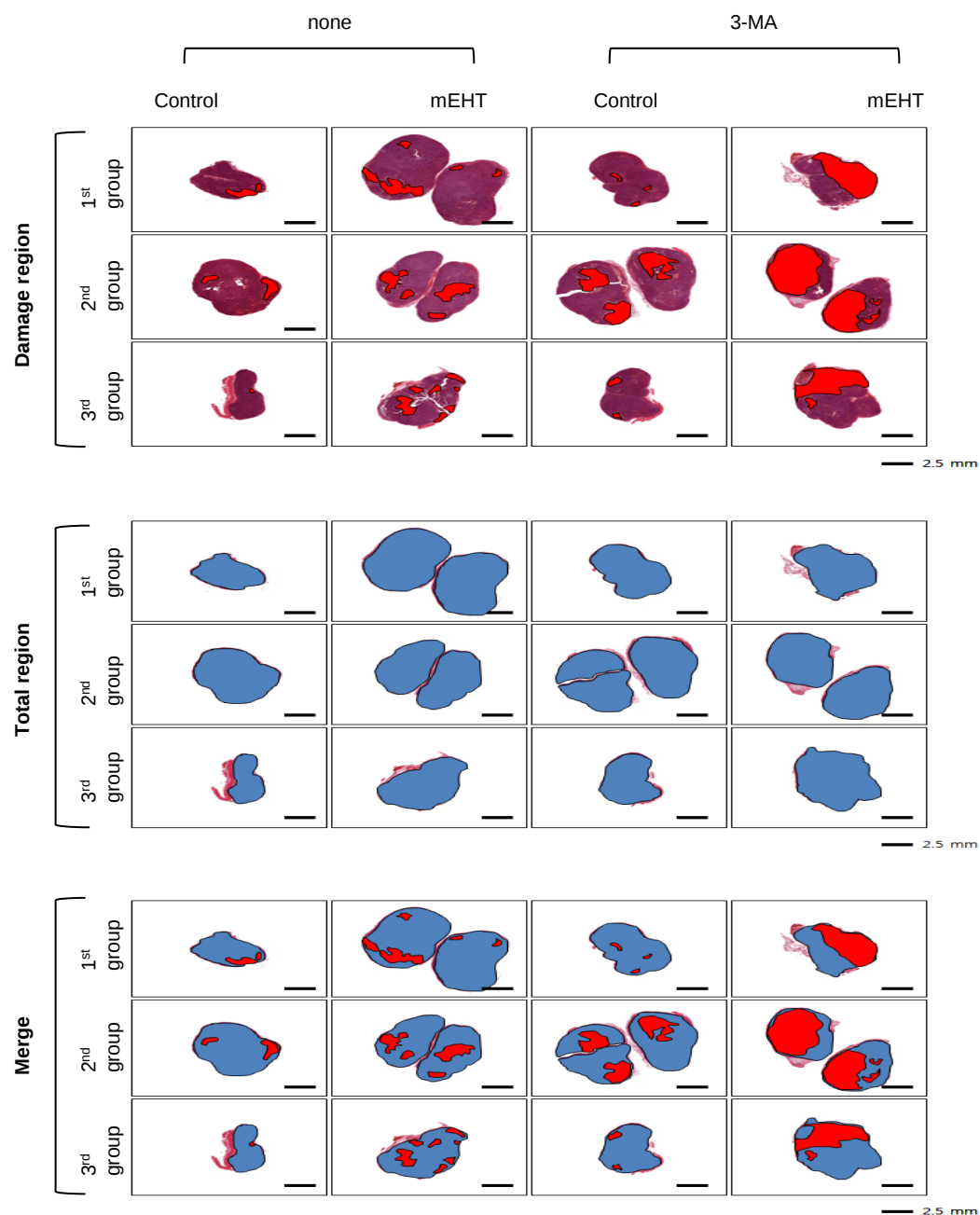


Figure S6. Subdivided images according to damaged tissue regions in haematoxylin and eosin (H&E) stained slides. Paraffin embedded xenograft tumor were sectioned at 10 μ m for H&E staining. H&E stained slide were scanned for high resolution image and were analyzed by pathologist. Pathologist have marked damaged region (Red) and whole tissue region (Blue) on H&E stained slides.

Table S1. Summary of mEHT results in PDTX-04 and PDTX-09.

PDTX number		n	Start day	Start Volume	End day	End volume	Variation of volume	p value
PDTX-04	Control	3	35	524.6 ± 86.5	44	940.9 ± 238.4	416.3 ± 171.5	0.1668 [#]
	mEHT	3		810.4 ± 201.2		673.1 ± 344.8	- 137.3 ± 279.7	
PDTX-09	Control	3	85	509.5 ± 101.8	104	771.1 ± 69.7	261.6 ± 62.3	0.0185 [*]
	mEHT	3		838.4 ± 145.4		804.8 ± 156.6	- 33.6 ± 45.1	

Results are the mean of growth variation (Δ tumor volume = tumor volume at start day – tumor volume at end day, tumor volume is mm³) ± S.E. n = 3. ([#], p > 0.05; ^{*}, p < 0.05)