

# Poly(d,l-lactide-co-glycolide)-polyethyleneimine/reduced nanographene oxide nanoparticles for chemo-phototherapy of cancer

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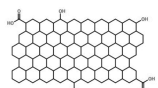
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## Background

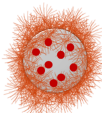
- Cancer illness such as triple-negative breast cancer lacks specific treatment targets which leads to poor treatment outcomes.<sup>[1]</sup>
- Photothermal therapy can synergistically work with chemotherapy.<sup>[2]</sup>
- Mild temperatures increase (39-45 °C) are able to induce cancer cell death and trigger drug release.<sup>[3]</sup>
- Graphene-based nanomaterials possess high near infrared absorption being ideal photothermal agents.<sup>[4]</sup>

**Reduced Graphene oxide**



- ✓ Higher NIR absorption
- ✓ Biocompatible

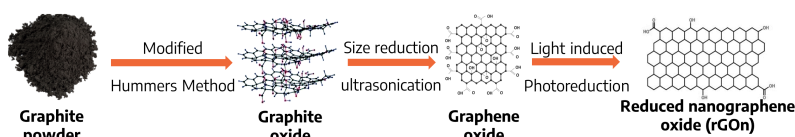
**PLGA-PEI Nanoparticles (NP)**



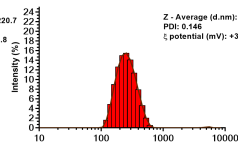
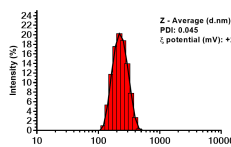
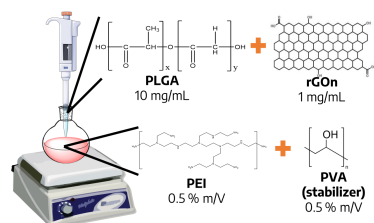
- ✓ High Loading capacity
- ✓ Enables surface moieties
- ✓ Biocompatible

- The copolymer of lactic glycolic acid (PLGA) is FDA approved, and allows high encapsulation yields for multiple drugs, such as the new one studied in this work - anticancer drug 1 (ACD-1).<sup>[5]</sup>
- Surface modification with polyethyleneimine (PEI) allows higher internalization and therapeutic effect.<sup>[5]</sup>

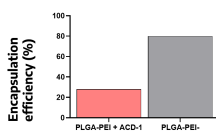
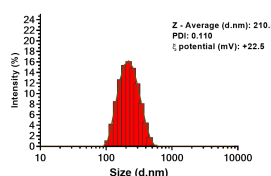
## Reduced Graphene Production



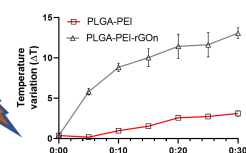
## Nanoparticle formulation



## Drug encapsulation



## Encapsulation of ACD-1



PLGA-PEI-rGO + ACD-1 NP

## NIR irradiation

## Conclusions

- PLGA-PEI-rGO particles were successfully produced, presenting sizes around 200 nm and being water stable.
- It was possible to encapsulate ACD-1 into those particles and rGO presence increased encapsulation efficiency by around 3.8-fold.
- rGO incorporation led to temperature increases above 13 °C, allowing to achieve values which permit their use for both triggered drug release and combined photothermal therapy.

## References

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