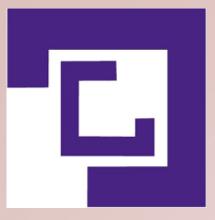
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# ELECTROCHEMICAL **CHARACTERIZATION OF GALLIC ACID**

**Department of Chemistry** 

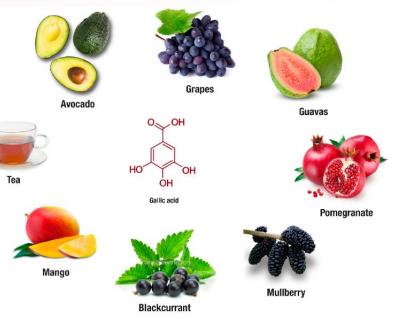
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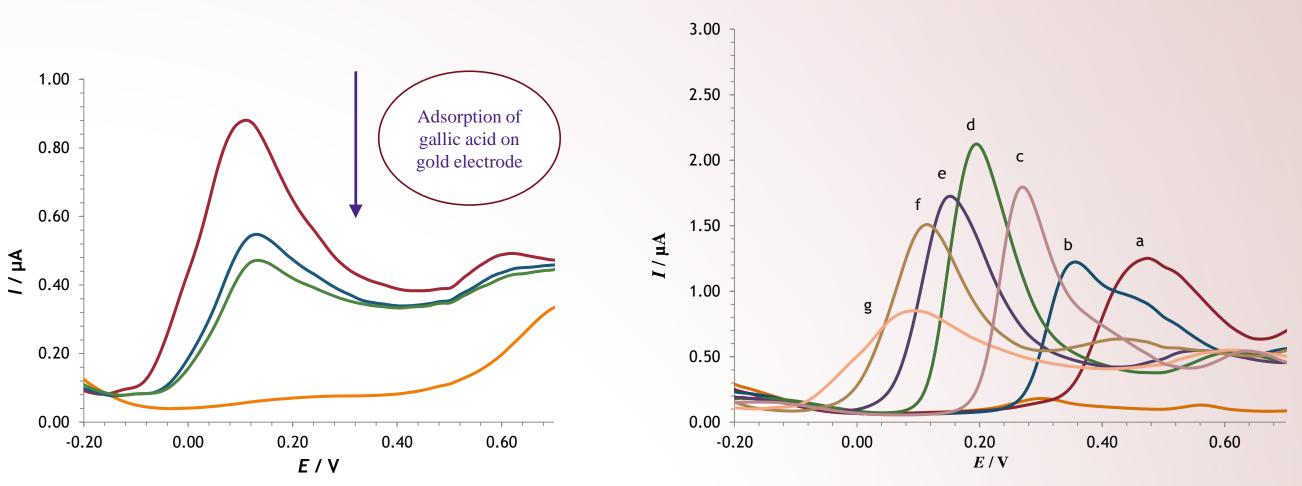
## **INTRODUCTION**

Gallic acid (3,4,5-trihydroxybenzoic acid) is a polyphenol, known for strong anticancer, antioxidant, anti-inflammatory and antimutagenic

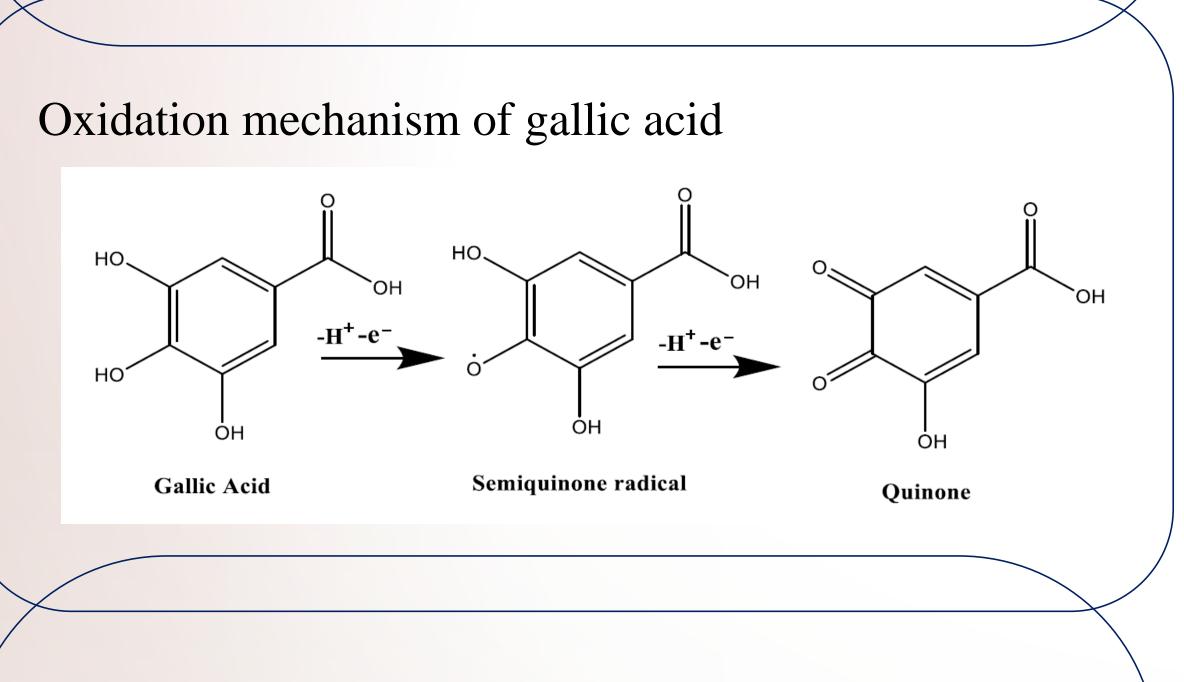
properties. Additional advantage of selective cytotoxicity for cancerous cells, makes gallic acid important additive in food to prevent cancer risks. It can be abudantly found, in different plants and foodstuff, especially in honey, gallnuts, mango,



### RESULTS



oak bark and pomegranate. The main goal of this study was to detect gallic acid in model systems and real samples (tea, liquer and wine) with differential pulse voltammetry.



#### **MATERIALS AND METHODS**

Fig. 1. Differential pulse voltamograms of gallic acid (c = 1x 10<sup>-3</sup> mol dm<sup>-3</sup>) in NaH<sub>2</sub>PO<sub>4</sub> ( $I_c$  = 0,1 mol dm<sup>-3</sup>):1<sup>st</sup> scan (—),  $2^{nd}$  scan (—) and  $3^{rd}$  scan (—). blank solution (—) NaH<sub>2</sub>PO<sub>4</sub>  $(I_c = 0, 1 \text{ mol dm}^{-3}).$ 

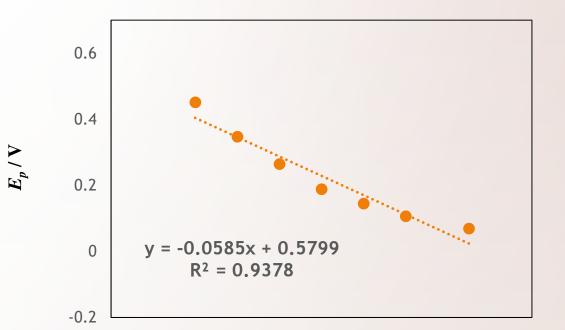
Fig. 2. Differential pulse voltamograms of gallic acid (c = $1 \ge 10^{-3} \mod \text{dm}^{-3}$  in NaH<sub>2</sub>PO<sub>4</sub> ( $I_c = 0,1 \mod \text{dm}^{-3}$ ) recorded at gold electrode as a function of pH (pH = (a) 3.0; (b) 4.0;(c) 5.0; (d) 6.0; (e) 7.0; (f) 8.0 and (g) 9.5).

**Table 1.** Values of peak potential  $(E_p)$  and peak current  $(I_p)$  of the first oxidation peak of gallic acid ( $c = 1 \ge 10^{-3} \mod \text{dm}^{-3}$ ) as a function of pH.

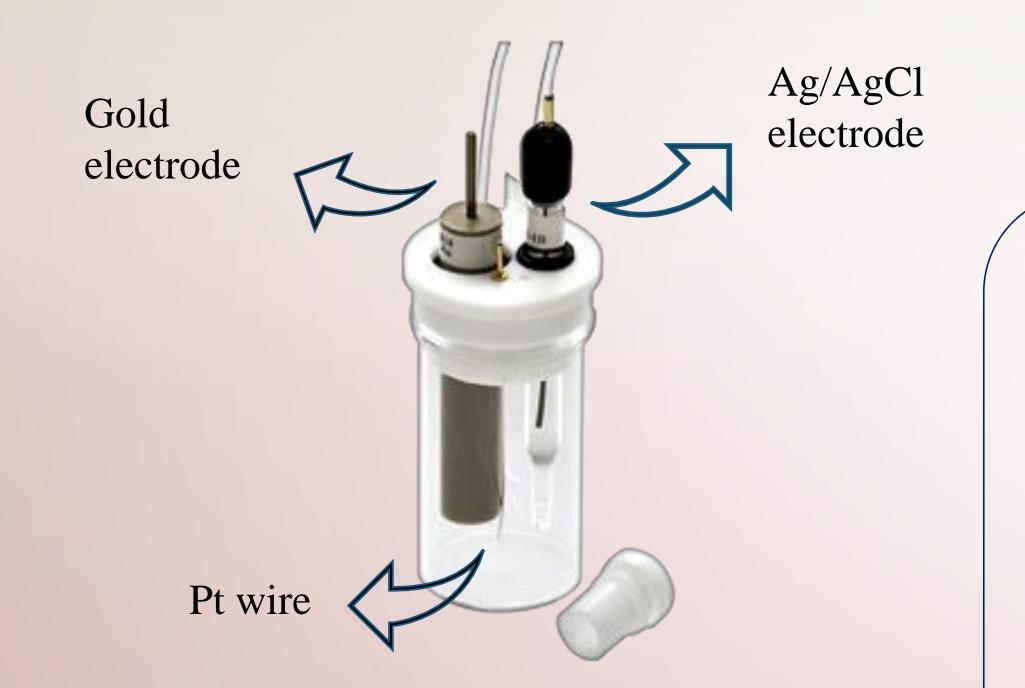
pН	$E_{\rm p}/{ m V}$	<i>I</i> <sub>p</sub> / μΑ
3.0	0.452	1.240
4.0	0.347	1.207
5.0	0.264	1.780
6.0	0.189	2.110
7.0	0.145	1.706
8.0	0.107	1.488
9.5	0.069	0.832

**Table 2.** Comparison of  $E_p$  and  $I_p$  values of gallic acid obtained with different working electrodes (gold (Au), platinum (Pt) and glassy carbon (GC)).

Woking electrode	$E_{ m p}$ / V	<i>I</i> <sub>p</sub> / μΑ
Pt	0.336	0.920
Au	0.187	1.749



- Electrochemical measurements were conducted in a three electrode voltammetric cell :
- 1. WORKING ELECTRODE: Au, Pt, glassy carbon (GC)
- 2. REFERENCE ELECTRODE: Ag/AgCl electrode
- 3. COUNTER ELECTRODE: platinum wire
- The system was purged with high purity argon, Ar5 ( $\phi_{Ar} =$ 99.999 %) before each measurement.
- Electrolyte: NaH<sub>2</sub>PO<sub>4</sub> ( $I_c = 0.1 \text{ mol dm}^{-3}$ )



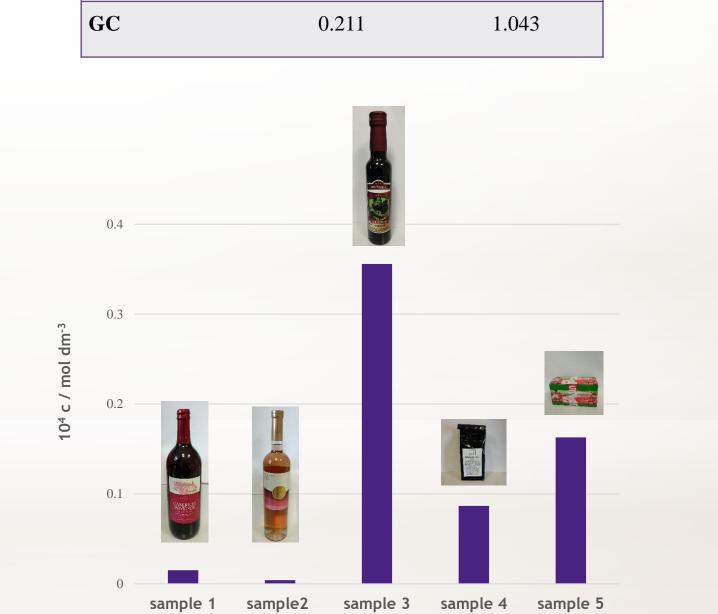


Fig. 5. Column graphs showing concentration of gallic acid in real samples.

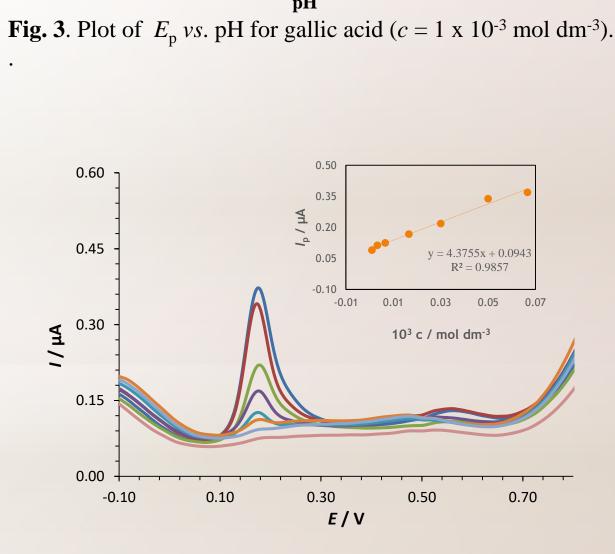


Fig. 4. Differential pulse voltammograms of gallic acid (c =1 -67  $\mu$ mol dm<sup>-3</sup>) in NaH<sub>2</sub>PO<sub>4</sub> ( $I_c$ = 0,1 mol dm<sup>-3</sup>) recorded at scan rate 5 mV/s. Inset: calibration curve for determination of gallic acid.

## CONCLUSIONS

- Two oxidation peaks of gallic acid were detected as well as adsorption of gallic acid oxidation product (quinone) on the gold electrode surface, oxidation of gallic acid was the most pronounced with Au working electrode
- The highest peak current of gallic acid was obtained around pH 6 while it decreased in more acidic and alkaline media,  $E_p$  vs. pH plot showed linearity with the slope 58.5 mV

