

Copper-Zinc Superoxide Dismutase (CuZn-SOD) Electrochemical Catalytic Amplification Sensing at Pt Ultramicroelectrodes

Annelis O. Sánchez-Álvarez^{1*}, J. Andres Melendez², Mariena Silvestry-Ramos³, Carlos R. Cabrera^{4*}

¹Department of Chemistry, University of Puerto Rico, Río Piedras Campus, San Juan, Puerto Rico 00931.

²College of Nanoscale Science and Engineering, SUNY Polytechnic Institute, Albany, New York 12203.

³Cornell Center for Material Research, Cornell University, Ithaca, New York 14853.

⁴Department of Chemistry and Biochemistry, University of Texas at El Paso, El Paso, Texas 79968.

***Corresponding author:** Annelis O. Sánchez-Álvarez, email: annelis.sanchez@upr.edu; Carlos R. Cabrera, email: crcabrerama@utep.edu

Received February 22nd, 2023; Accepted July 9th, 2023.

DOI for the article: <http://dx.doi.org/10.29356/jmcs.v67i4.1963>

Supplementary Information

Catalytic amplification numerical data analysis

Table SI 1. Some examples of peaks analysis by catalytic amplification.

I_{initial} (A)	I_{final} (A)	ΔI	r (cm) distance from active site	r (nm) distance from active site
3.08E-11	3.11E-11	2.80E-13	8.67E-08	0.87
2.14E-11	2.18E-11	3.90E-13	1.21E-07	1.21
1.86E-11	1.88E-11	2.50E-13	7.74E-08	0.77
2.33E-11	2.36E-11	2.30E-13	7.12E-08	0.71
2.73E-11	2.75E-11	2.10E-13	6.50E-08	0.65
1.79E-11	1.82E-11	3.40E-13	1.05E-07	1.05
2.03E-11	2.06E-11	2.80E-13	8.67E-08	0.87

$$r = \Delta I / 4\pi(\ln 2)nFDC \quad (1)$$

Where

n = # of e- transferred during the process

F = Faraday's constant

D = diffusion coefficient of O_2 at 20 °C in a ~100 mM salt solution [1]

C = concentration of O_2 in saturation for aqueous solution [2]

Calculation of theoretical diffusion coefficient using Stokes-Einstein equation

Stokes-Einstein approximation is described by the following equation [3]:

$$D_E = \frac{k_B T}{6\pi\eta r} \quad (2)$$

where k_B is Boltzmann's constant, T is absolute temperature (298.15 K) for these set of experiments, η is the solution's viscosity, and r is the nanoparticle's radius (2.1×10^{-9} m used as theoretical radius). The value obtained using this calculation was 1.24×10^{-6} cm²/s.

Frequency of collisions and experimental diffusion coefficient

The experimental frequency of collisions was calculated using the number of peaks per 300 s ratio. These values were plotted against the enzyme concentration in solution. Since frequency of collision due to diffusion (f) can be described as follows[4]:

$$f = 4D_E C_E r_{elec} N_A \quad (3)$$

the experimental diffusion coefficient was calculated using the experimental frequency at known concentration:

$$\text{i.e. } (C_E = 0.9 \text{ fM}); D_E = \frac{f}{4C_E r_{elec} N_A} = \frac{.00333 \text{ s}^{-1}}{(4)(9E^{-19} \frac{\text{mol}}{\text{cm}^3})(6.02E^{23} \text{ mol}^{-1})(5E^{-4} \text{ cm})} = 3.08 \times 10^{-6} \text{ cm}^2/\text{s}$$

The magnitude of the experimental diffusion coefficient was 1×10^{-6} cm²/s regardless of the concentration used for the positive slope lineal area.

References

1. Jamnongwong, M.; Loubiere, K.; Dietrich, N.; Hébrard, G. *Chem. Eng. J.* **2010**, 165, 758-768.
2. Robinson, J.; Cooper, J. M. *Anal. Biochem.* **1970**, 33, 390-399.

3. Einstein, A. *Annalen der Physik.* **1905**, 322, 891-921.
4. a) Kim, B.-K.; Kim, J.; Bard, A. J. *J. Am. Chem. Soc.* **2015**, 137, 2343-2349; b) Sekretaryova, A. N.; Vagin, M. Y.; Turner, A. P. F.; Eriksson, M. *Procedia Technol.* **2017**, 27, 238-239; Sekretaryova, cA. N.; Vagin, M. Y.; Turner, A. P. F.; Eriksson, M. *J. Am. Chem. Soc.* **2016**, 138, 2504-2507.