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# Performance of air-cathode stacked microbial fuel cells systems for wastewater treatment and electricity production

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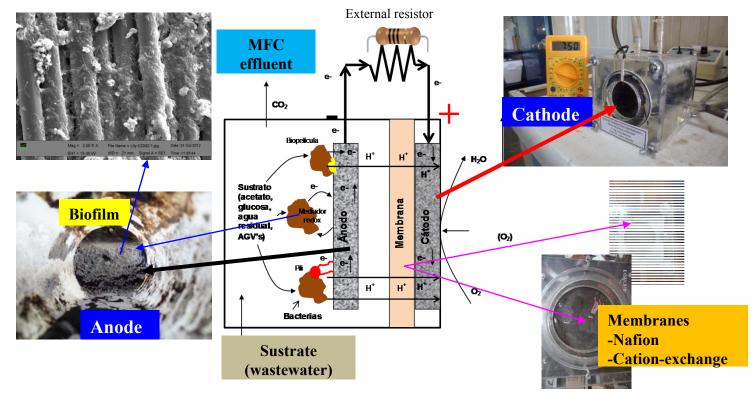




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The MFCs are bio-electrochemical systems, capable of generating electricity from the oxidation of organic matter from wastewater treatment and simultaneously the contaminant removal.



\* Application for wide locations and diverse fuels

Advantages of MFC over current energy generating technologies from organics

\* High conversion efficiency

- \* Ambient temperature
- \* No gas treatment
- \* No energy input for aeration



# **Principies of MFC**



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• Biochemical degradation – microorganism growth : Oxidation-reduction

Electrochemistry, Catalyst reaction

Mass transport, Mixing of substrate

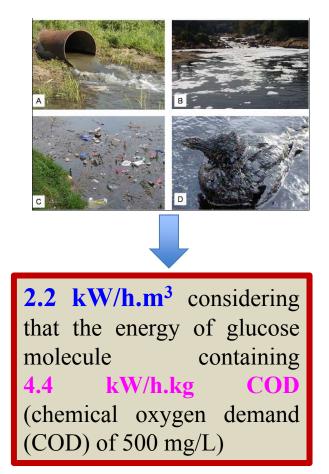
• @ anode, acetate (substrate) is oxidized by bacteria (catalyst), mixed culture.

$$CH_3COO^- + 4 H_2O \rightarrow 2 HCO_3^- + 9H^+ + 8e^-$$
 (E<sub>an</sub> = -0.296 V)

**Electrons** produced are transferred to the anode by mediator/shuttles, direct membrane, or nano-wire.

- Electrons flow to cathode through a conductive material containing a resistor or operated under a load to produce electricity.
- **Protons** diffuse from anode to cathode through cation exchange membrane.
- (a) cathode, the electrons are combined with proton and **oxygen** or chemical oxidizer. Diverse bacterial community is working as a catalyst.

$$O_2 + 4H^+ + 4e^- \rightarrow 2H_2O$$
 (E<sub>cat</sub> = 0.805 V)





## **Voltage productions by MFCs**

Cathode

 $\mathrm{E}_{\mathrm{cat}}$ 



#### Single chamber air-cathode MFC

- Standard electrode potential, at SC (25 °C, 1 atm, 1 M) = zero, relative to normal hydrogen electrode (NHE)
- Maximum attainable cell voltage can be calculated by,

$$E_{emf} = E_{cat} - E_{an}$$

According to Nernst equation.

$$E_{emf} = E_{emf}^{0} - \frac{RT}{nF} In \left( \frac{[C]^{v_{C}}[D]^{v_{D}}}{[A]^{v_{A}}[B]^{v_{B}}} \right)$$

Acetate oxidized at anode, oxygen used as e-acceptor at cathode

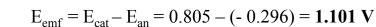
Anode 
$$2 \operatorname{HCO}_3^- + 9 \operatorname{H}^+ + 8 \operatorname{e}^- \rightarrow \operatorname{CH3COO}^- + 4 \operatorname{H}_2 \operatorname{O}$$

 $O_2 + 4H^+ + 4e^- \rightarrow 2H_2O$ 

E<sub>an</sub> = 
$$E_{an}^{0}$$
 - RT/8F ln ([CH<sub>3</sub>COO-]/[HCO3-]<sup>2</sup>[H+]<sup>9</sup>) = -0.296 V

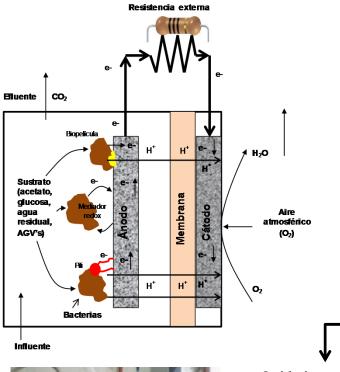
$$= E_{eat}^{0} - RT/4F \ln (1/pO_{2}[H+]^{4}) = 0.805 V$$

Oxidation-reduction potentials



#### Maximum MFC voltage

Theoretical = **1.1 V** Open circuit mode (OCV)) = **0.6-0.8 V(without current)** Real voltage during current generation < **0.62 V** 



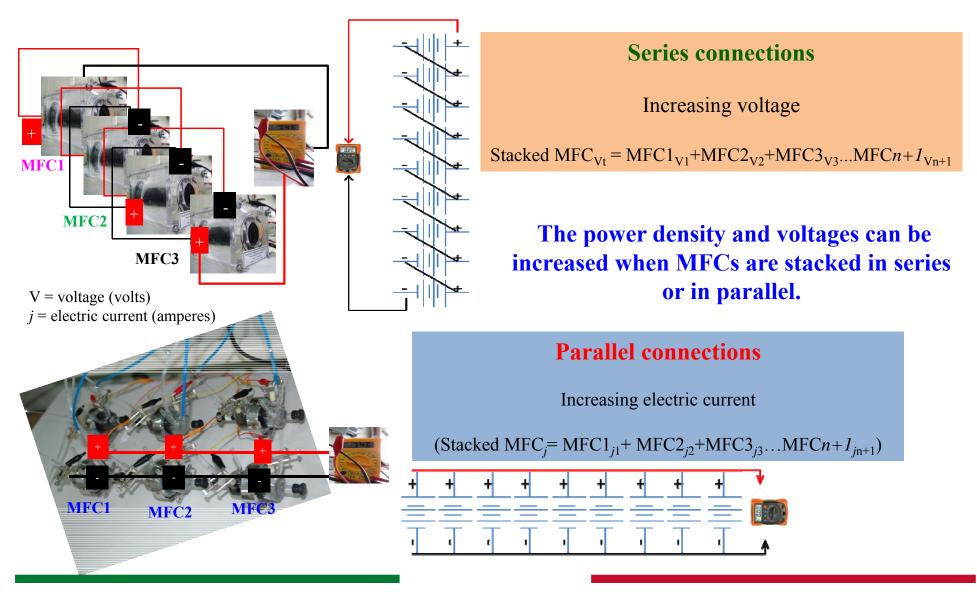


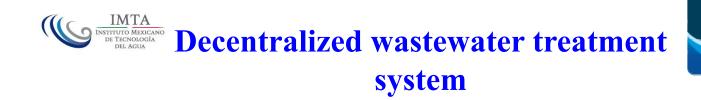
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## **Stacked MFC (Multi-electrode MFCs)**

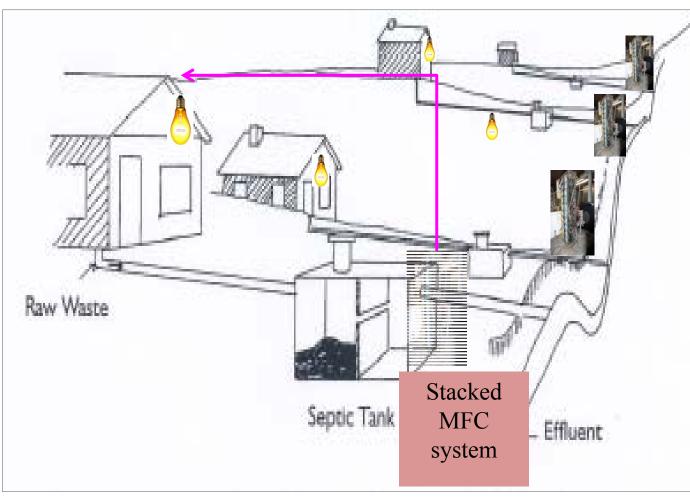


A single cell delivering about 0.2-0.8 V (too low for most applications). Just like batteries, individual cells are stacked to achieve a higher voltage and power. This assembly of cells is called a cell stack, multi-electrode or just a stack.





#### MFCs as alternative for wastewater treatment.



New Process-Microbial Electrochemical

Technologies

\* Removal organic matter

\* Electricity direct

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### **Objective**

The main objective of this study was to evaluate the performance of two air-cathode stacked MFC systems at different HRT (3, 1 and 0.5 d) during wastewater treatment and electricity production.



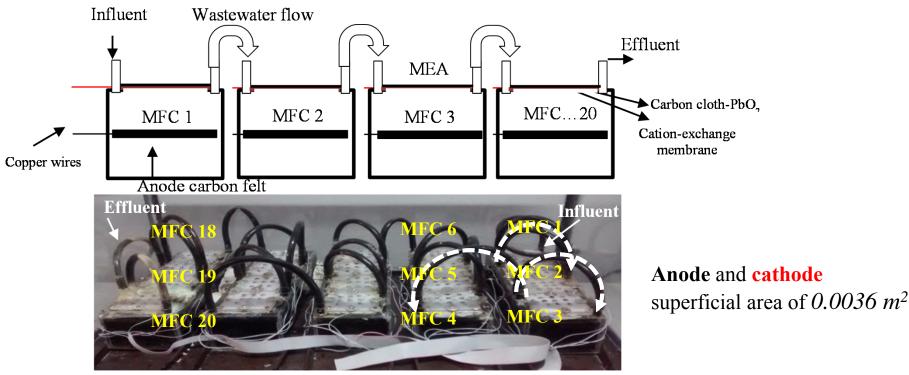


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#### Architecture of stacked MFCs

#### Stacked MFC 1 (un-shared reactor)-shared anolyte



- 20 individual MFC unit (800 mL MFC); Total volume 16 L.
- Single chamber air-cathode MFC
- Stacked MFC 1 was fed in continuous flow cascade mode (flow was transported through of the each MFC compartment

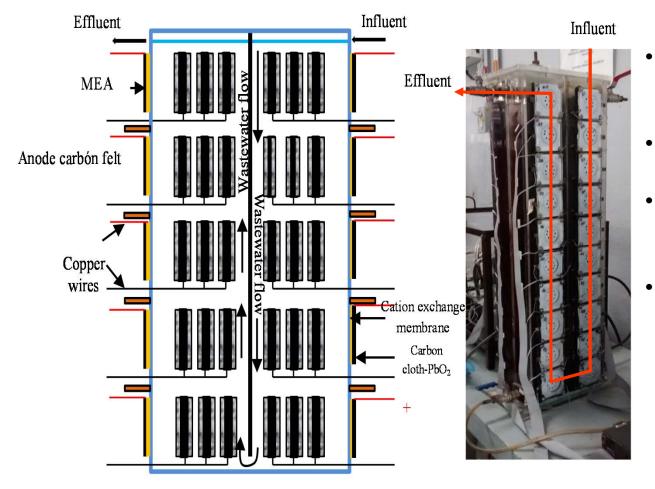




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#### Stacked MFC 2 (shared reactor)-shared anolyte



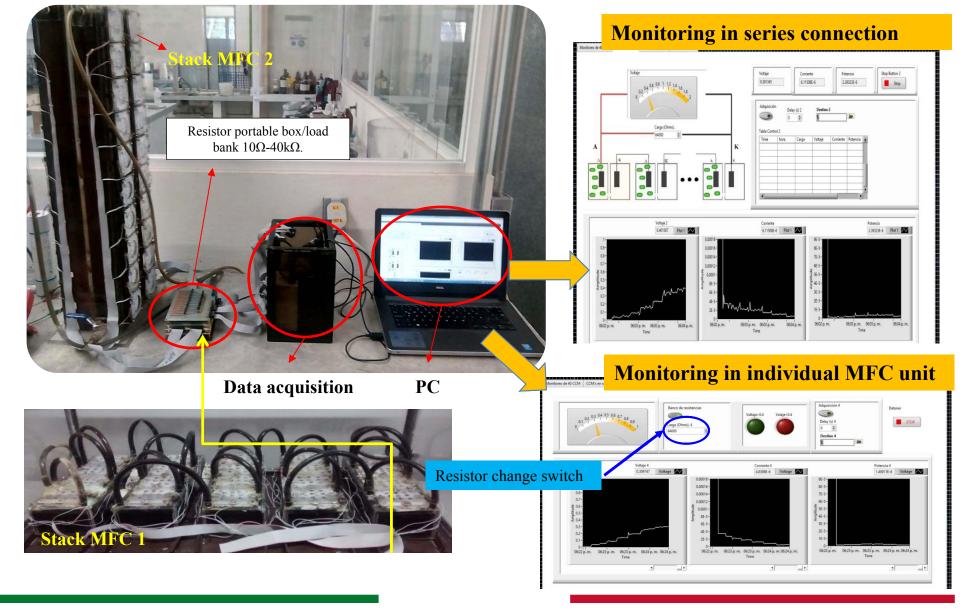
- 40 MFC unit into shared reactor.
- Total volume 16 L.
- Single chamber air-cathode MFC.
- Stacked MFC 2 was fed in continuous flow (not separator between in each cell was used).





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#### Table 1. Operational conditions

	Open Circuit Mode	Closed Circuit Mode (1,000 Ω)
Biomass aclimatation	HRT 10 d	-
HRT (d)	3, 1 and 0.5	3, 1 and 0.5

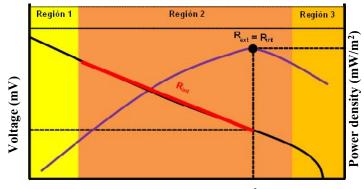
#### INDIVIDUAL MFC UNIT SERIES CONNECTION



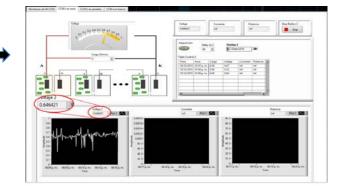
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#### **Polarization curves**







COD, TN, and TP were measured using standard methods (APHA, 2005).

$$P = V * I = \frac{V^2}{R_{ext}} = I^2 * R_{ext}$$

$$P^* = \frac{V^2}{aR_{ext}} = \frac{P}{a}$$
$$I = \frac{dq}{dt} = \frac{V}{R_{ext}}$$
$$J = \frac{q}{at} = \frac{1}{a} \left(\frac{q}{t}\right) = \frac{I}{a}$$





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# Characteristic of municipal wastewater

 $COD = 209 \pm 41 \text{ mg/L}.$ 

Total Nitrogen (TN) =  $38 \pm 11 \text{ mg/L}$ .

Total Phosphorus (TP) =  $15 \pm 3$  mg/L.

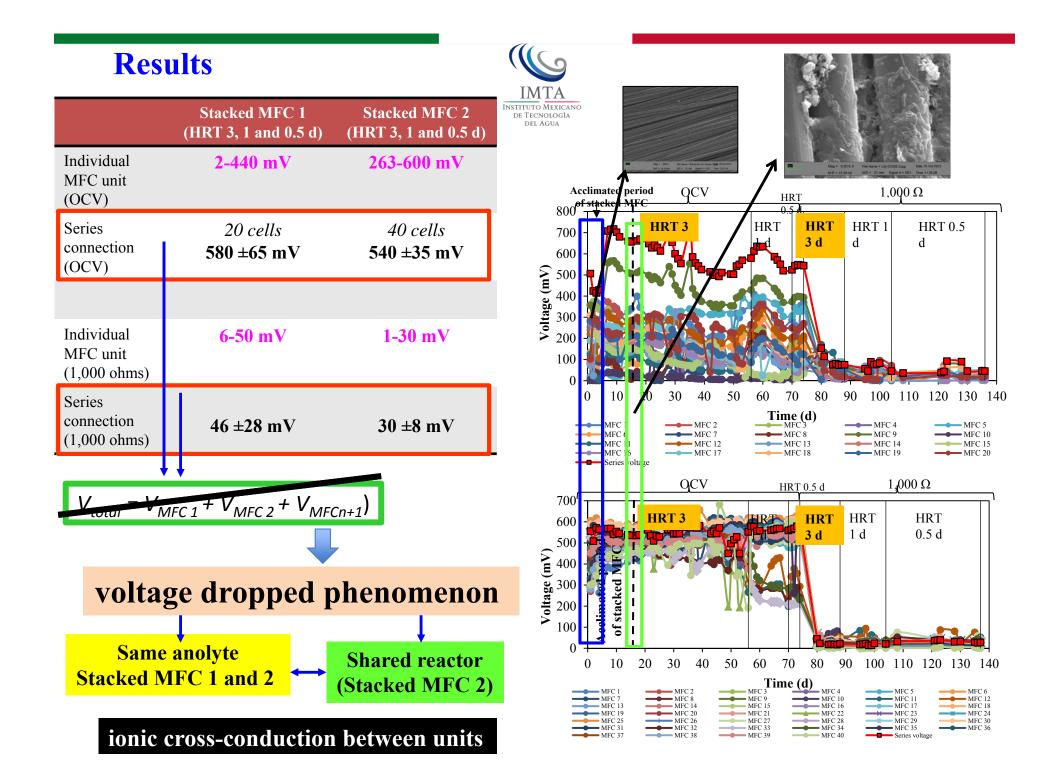
pH = 6.9-7.5

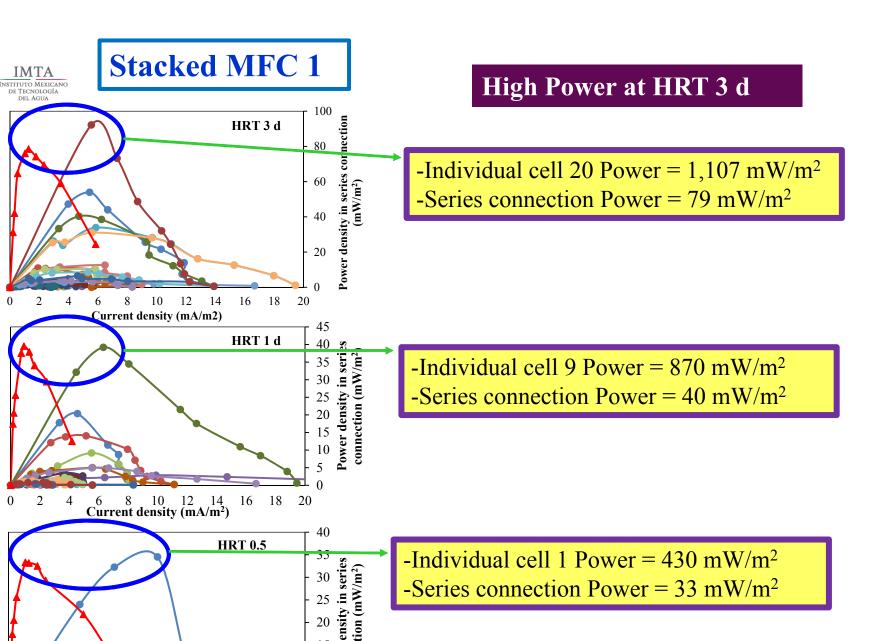
Total Suspended Solids (TSS) = 200 mg/L

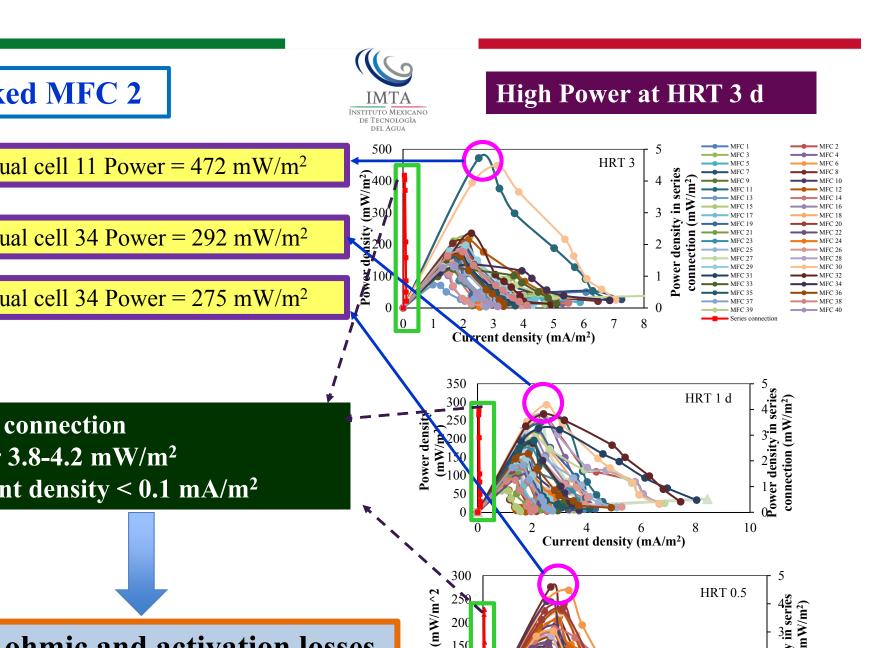
**Residential housing** 





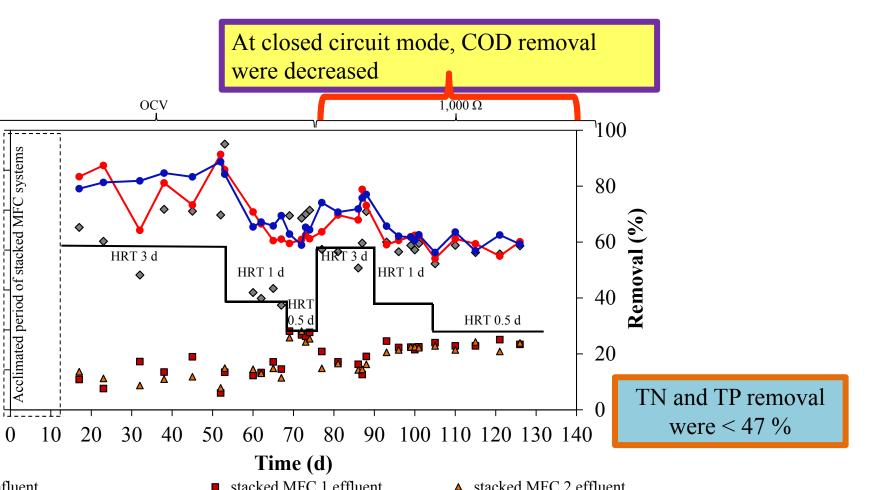








# Removal COD from Stacked MFC 1 and 2 at different HRT (3, 1 and 0.5 d





#### Conclusions

- The two stacked MFC systems tested were not effective for power production in series connection under OCV and closed circuit mode.
- Voltages dropped (voltages reversal) in the two systems were generated due to the architecture of the systems (shared reactor) and the same anolyte in all MFC units.
- The maximum power density in series connection of the stacked MFC 1 was 79  $\pm 0.65 \text{ mW/m}^2$  (current density of  $1.3 \pm 0.4 \text{ mA/m}^2$ ). For the individual MFC unit (no-series connection), the maximum power density was  $1,106 \pm 1.2 \text{ mW/m}^2$  (current density of 5.5 0.6 mA/m<sup>2</sup>) at HRT of 3 d.
- Power production in stacked MFC 2 ( $4.2 \pm 0.6 \text{ mW/m}^2$ ) and the current density ( $0.04 \pm 0.006 \text{ mA/m}^2$ ) were lower compared with the power generated from stacked MFC 1.
- The results showed that the COD removal(up to 84%) were increased when the HRT were increased from 0.5 to 3 d.







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