



6rd DRD1 Meeting

Varsaw, Oct. 2025

Unveiling Surface Chemistry and Aging Pathways of GEMs through In-Situ Surface Spectroscopy

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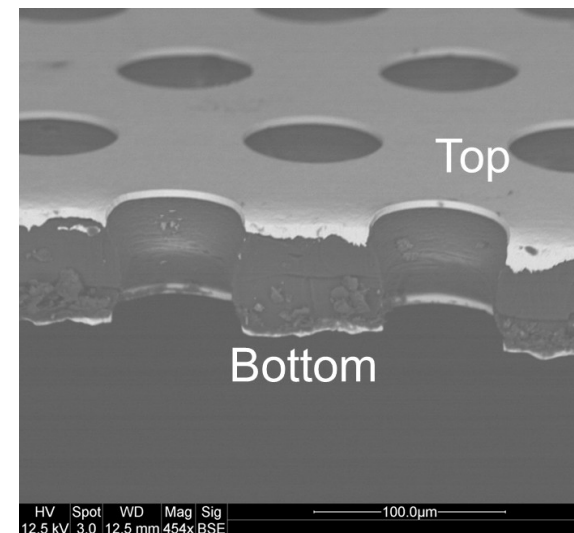
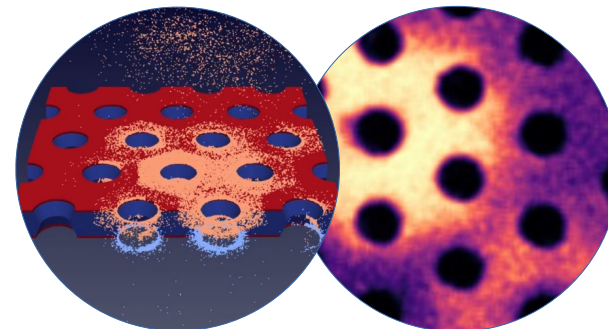
Maria Do Carmo M. Alves, Jonder Moraes

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High Energy Physics and Instrumentation Center at USP

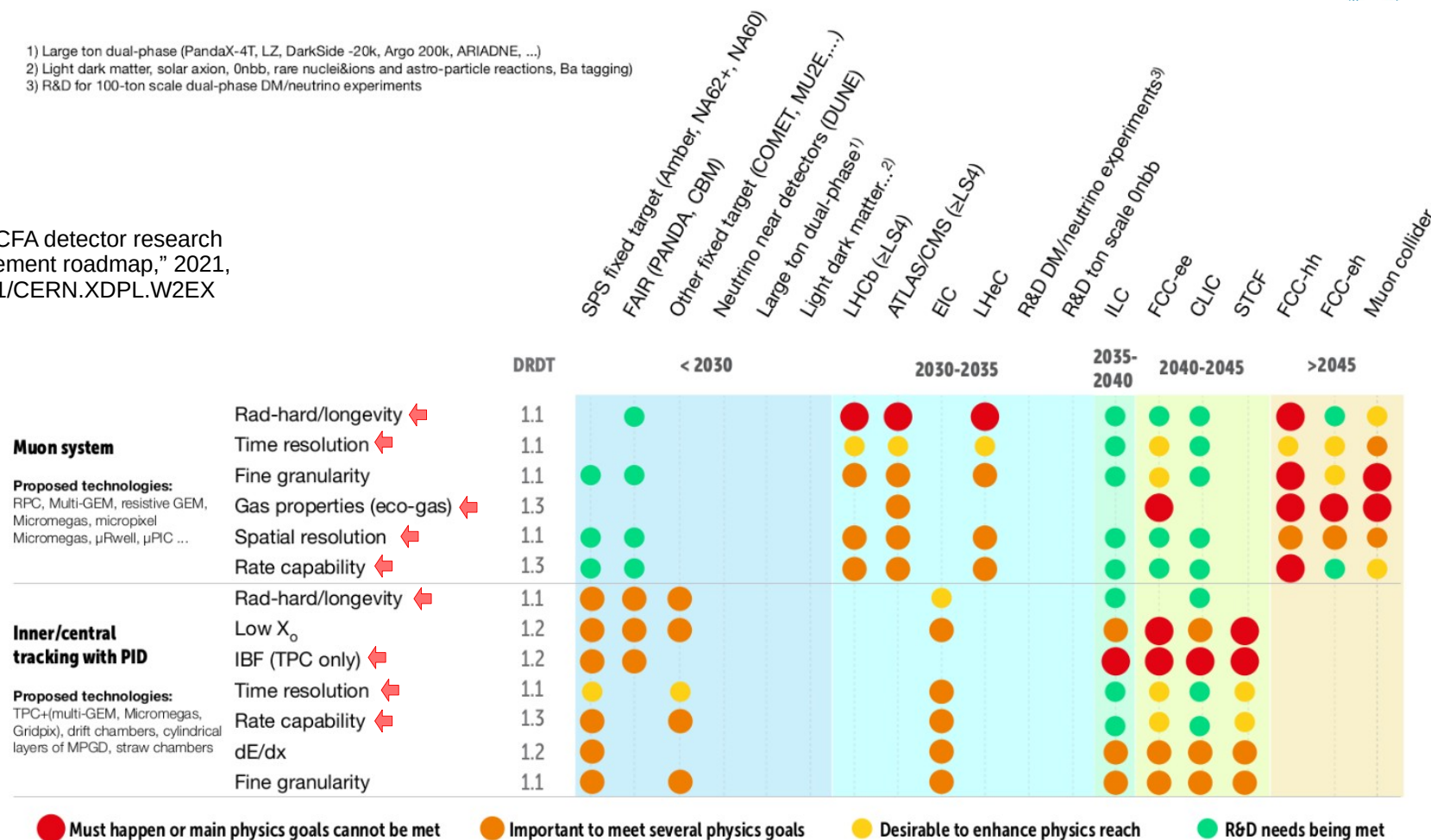
- **Reliability & Stability:** Ensure long-term gain uniformity and tracking precision.
- **Predictive Maintenance:** Detect early degradation to prevent failures and reduce costs.
- **Radiation & Charge Effects:** Understand how ionization and charge buildup alter GEM performance.
- **Surface Chemistry:** Investigate gas–material interactions, contamination, and polymerization on GEM surfaces.
- **Ion Feedback & Discharges:** Mitigate micro-discharges and feedback under high-rate operation.
- **Design Optimization:** Guide material selection, cleaning, and fabrication for next-generation GEMs.

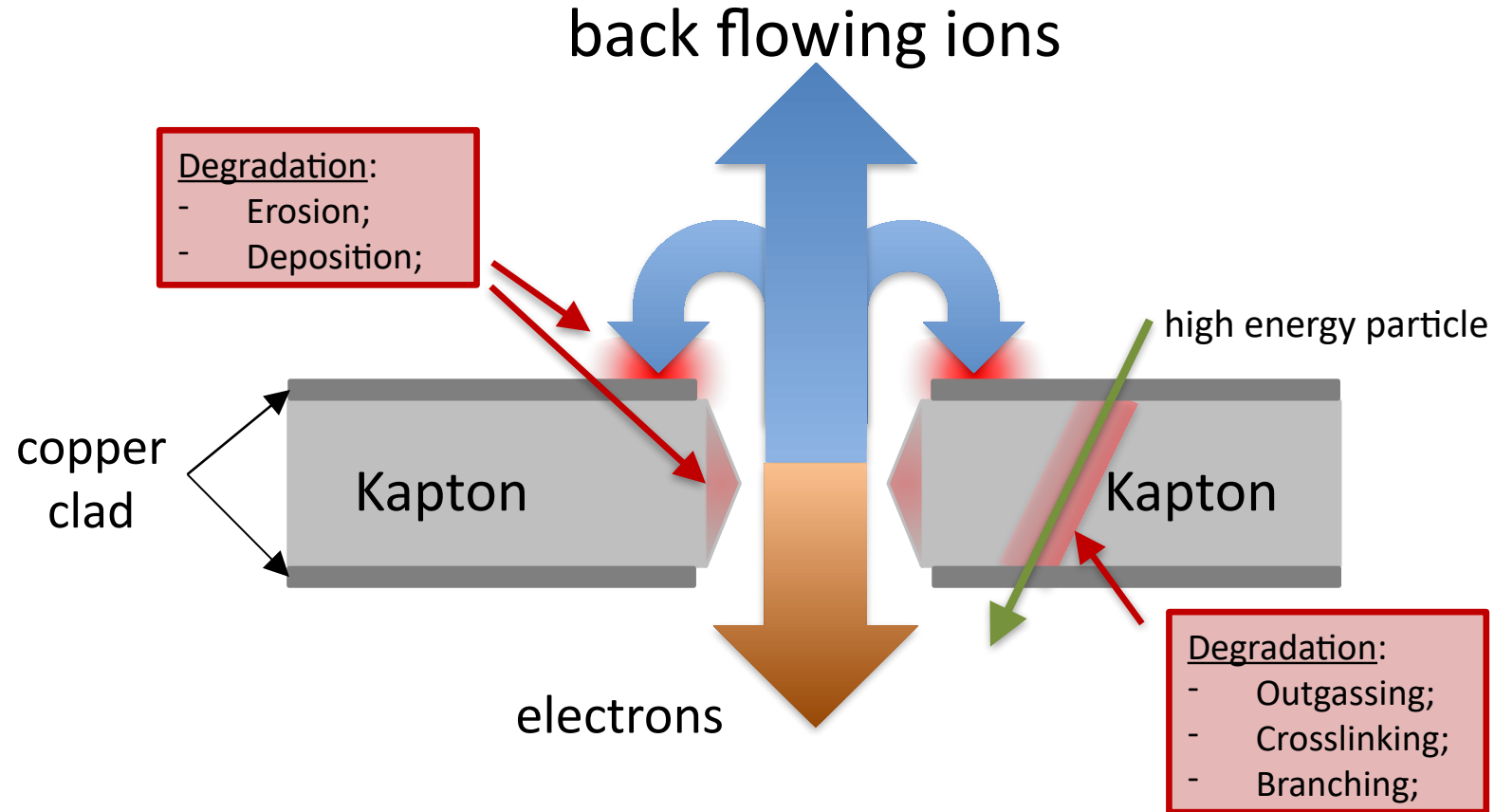


Aging and degradation models: why studying it?

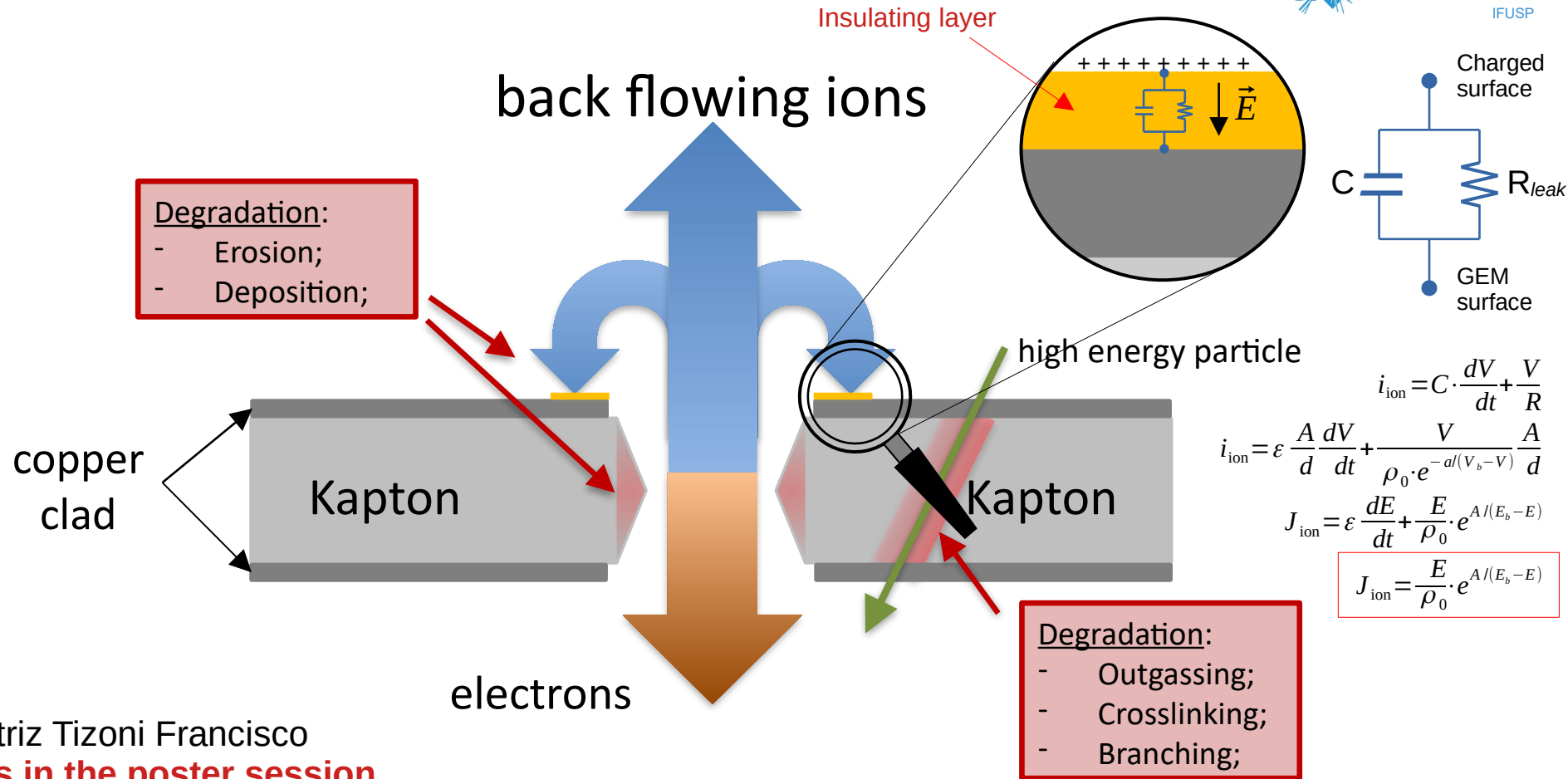
- 1) Large ton dual-phase (PandaX-4T, LZ, DarkSide -20k, Argo 200k, ARIADNE, ...)
- 2) Light dark matter, solar axion, Onbb, rare nuclei&ions and astro-particle reactions, Ba tagging)
- 3) R&D for 100-ton scale dual-phase DM/neutrino experiments

“The 2021 ECFA detector research and development roadmap,” 2021, doi: 10.17181/CERN.XDPL.W2EX





Aging and degradation models: the Malter effect

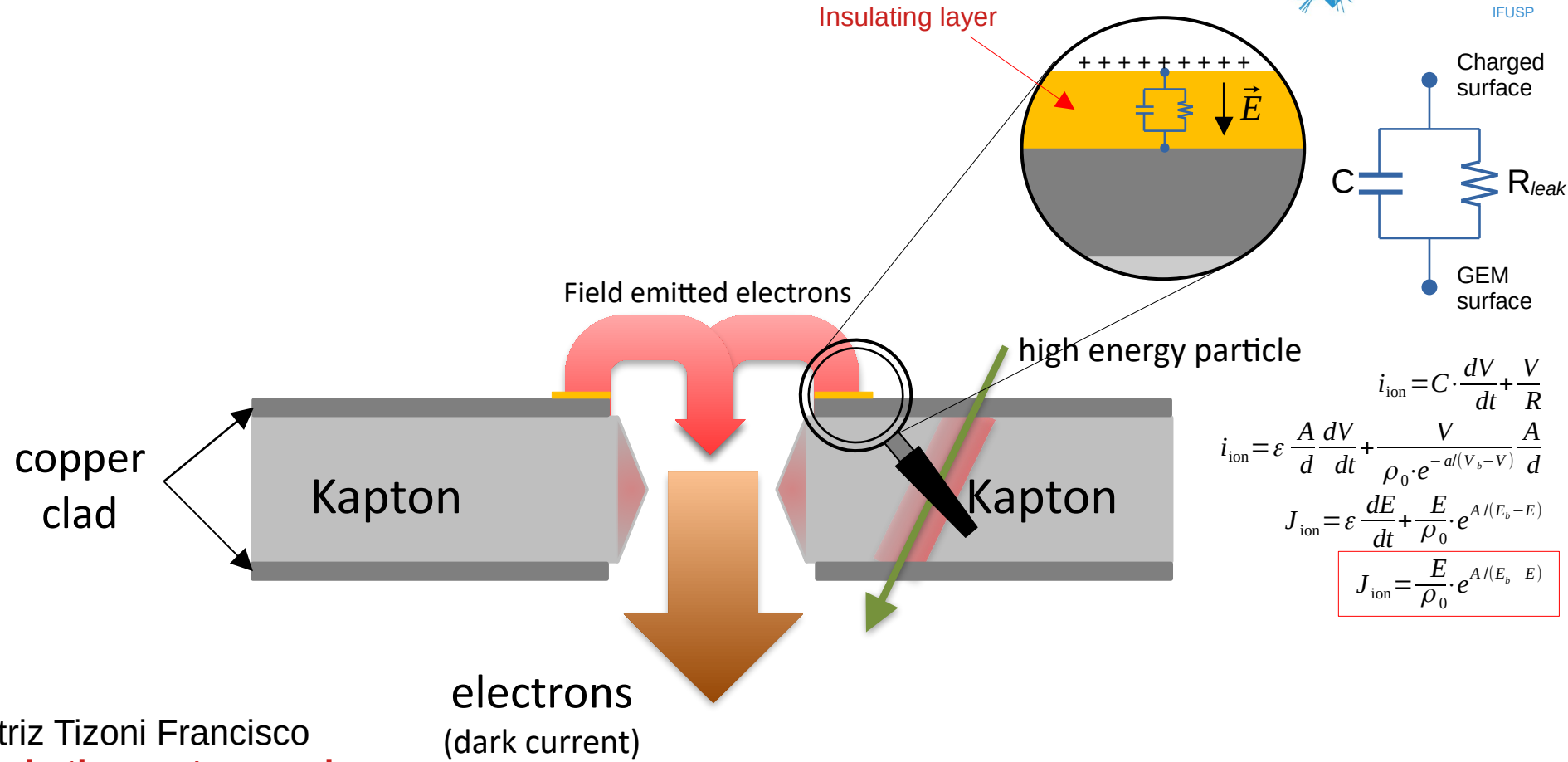


Bruna Beatriz Tizoni Francisco

See details in the poster session

“Understanding the dynamics of the Malter effect in GEMs through computer simulations”

Aging and degradation models: the Malter effect

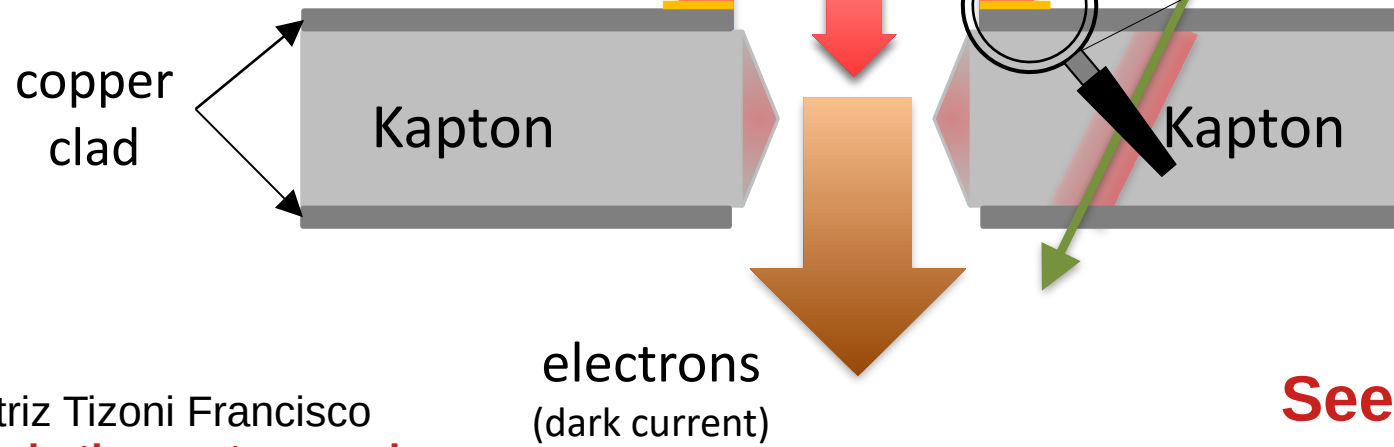
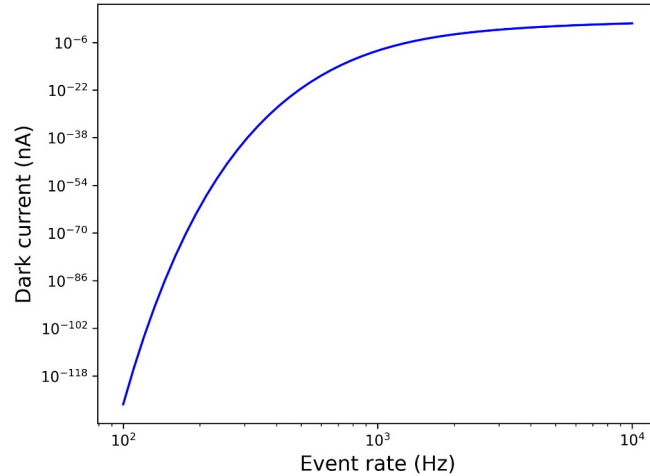


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Aging and degradation models: the Malter effect



$$i_{\text{ion}} = C \cdot \frac{dV}{dt} + \frac{V}{R}$$

$$i_{\text{ion}} = \varepsilon \frac{A}{d} \frac{dV}{dt} + \frac{V}{\rho_0 \cdot e^{-a/(V_b - V)}} \frac{A}{d}$$

$$J_{\text{ion}} = \varepsilon \frac{dE}{dt} + \frac{E}{\rho_0} \cdot e^{A/(E_b - E)}$$

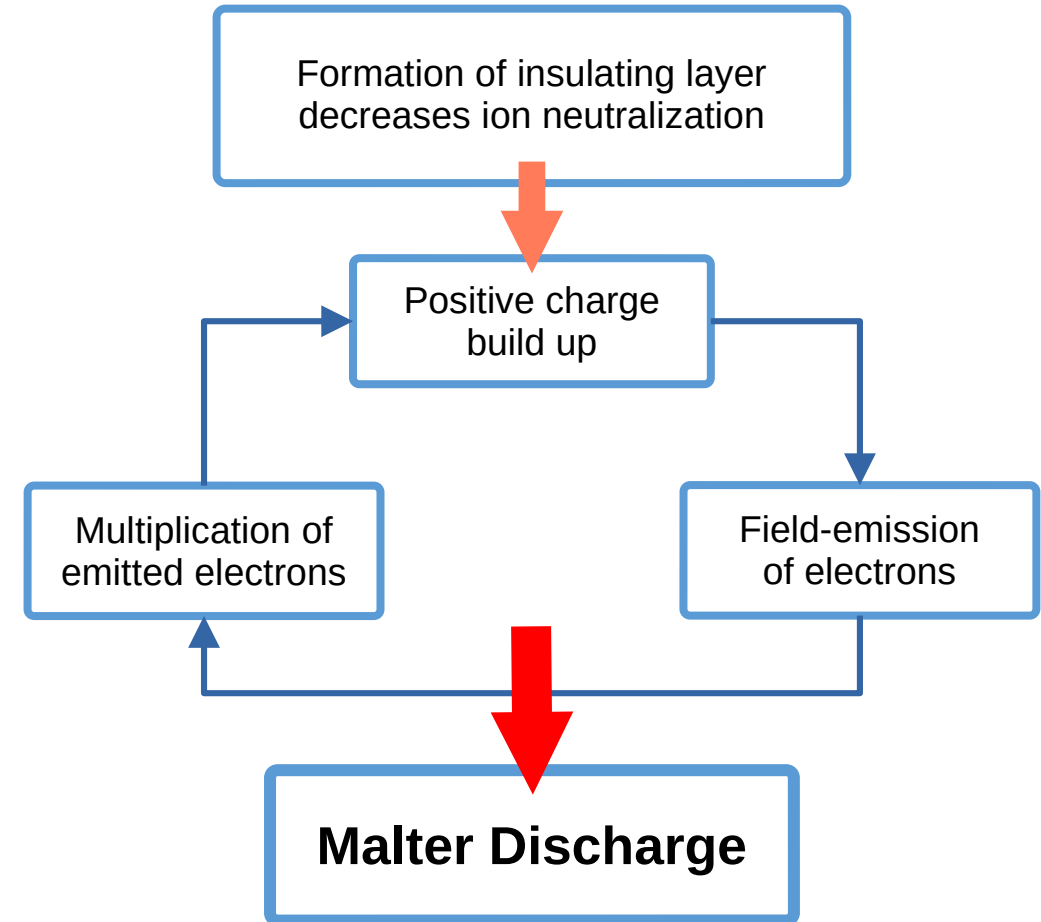
$$J_{\text{ion}} = \frac{E}{\rho_0} \cdot e^{A/(E_b - E)}$$

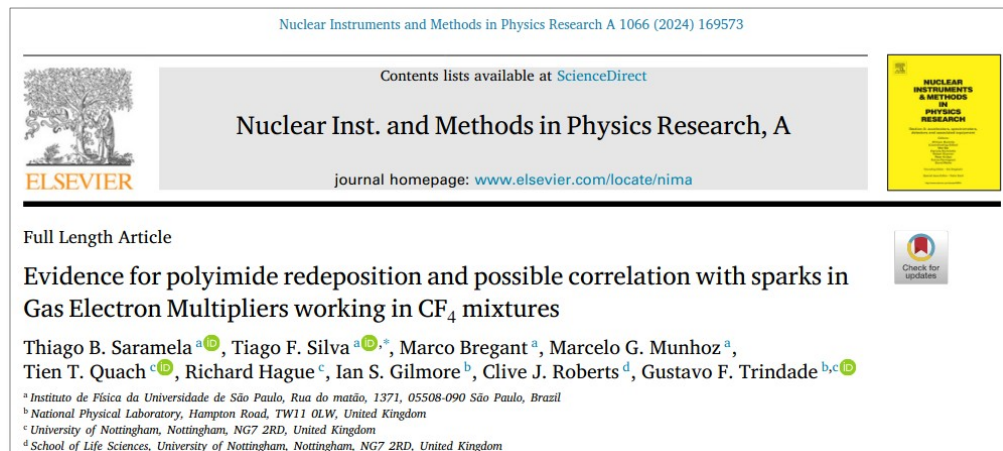
See poster 21!

Bruna Beatriz Tizoni Francisco
See details in the poster session

“Understanding the dynamics of the Malter effect in GEMs through computer simulations”

- **Discharge Behavior:** Films modify surface conductivity, increasing the risk of micro-discharges.
- **Surface Chemistry:** Understand oxidation, adsorption, and polymerization processes on copper.
- **Aging Mechanisms:** Identify how radiation, gas composition, and impurities drive film growth.
- **Material Optimization:** Guide cleaning, treatment, and coating methods to control surface passivation.



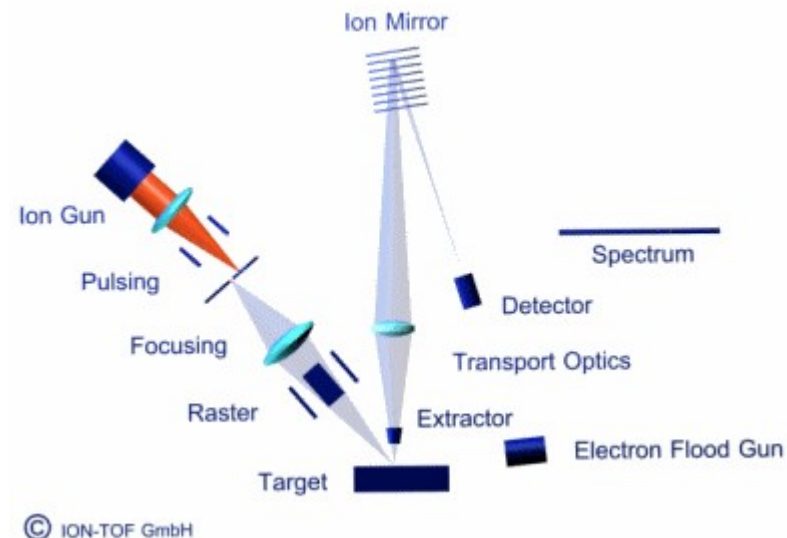


ToF-SIMS is a ultra sensitive surface analysis technique.

The University of Nottingham equipment

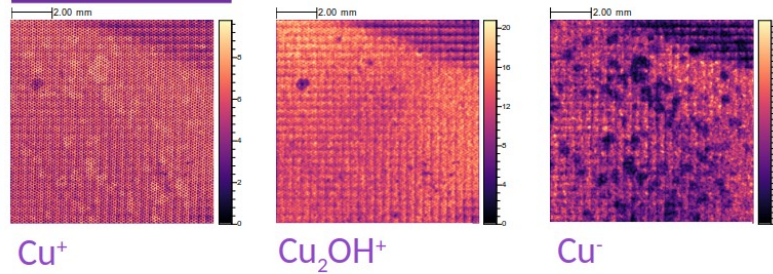
• Iontof TOF.SIMS IV

- Liquid metal primary ion source (Bi_n^+)
- Gas Cluster ion Beam source (Ar_n^+)
- Thermal sputtering ion Source (Cs^+)
- Single stage reflectron ToF analyser
- Nominal mass resolving power m/dm @ 29 u: 10,000

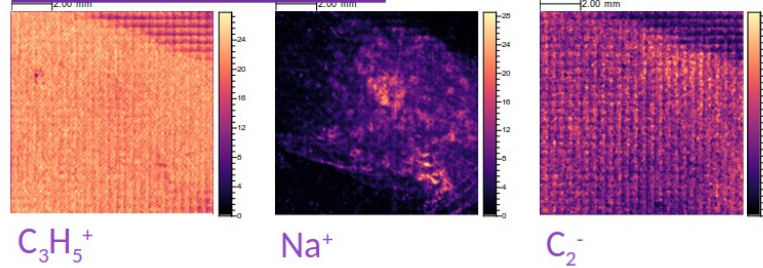


ToF-SIMS as a surface analysis technique

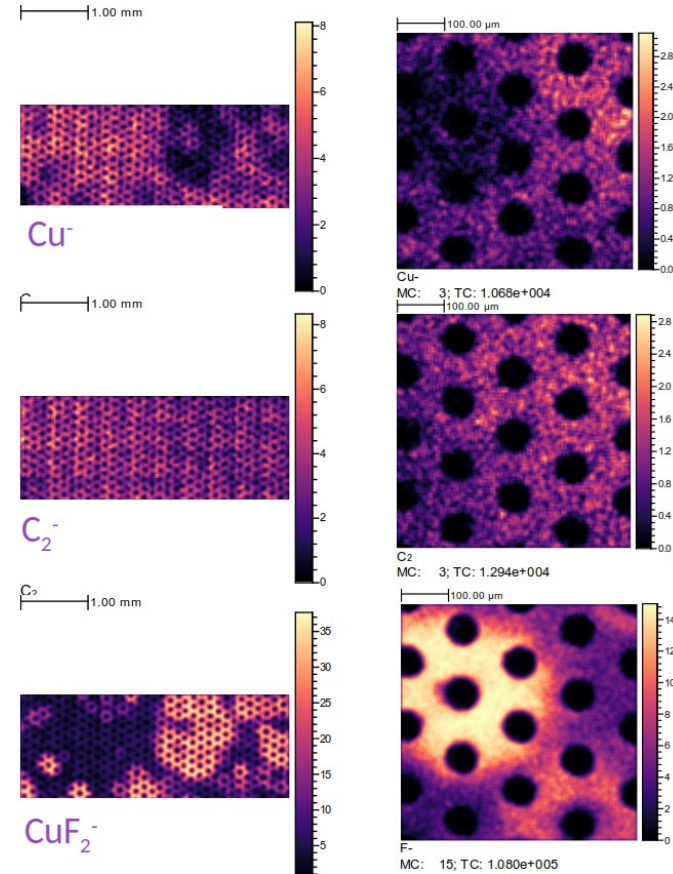
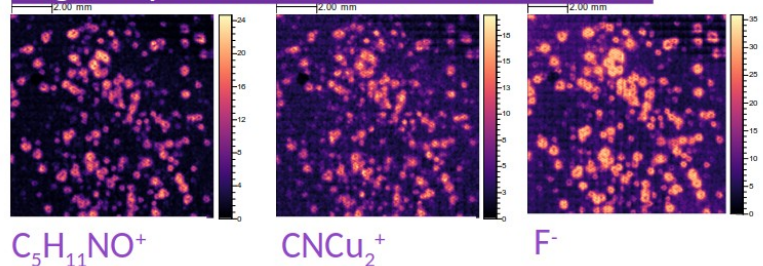
Cu oxide / Cu



Surface contamination

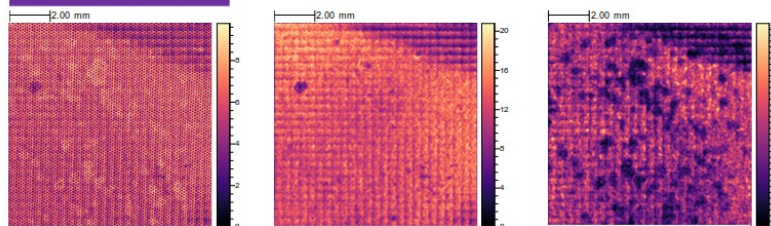


Organic pattern adsorbed onto surface



ToF-SIMS as a surface analysis technique

Cu oxide / Cu

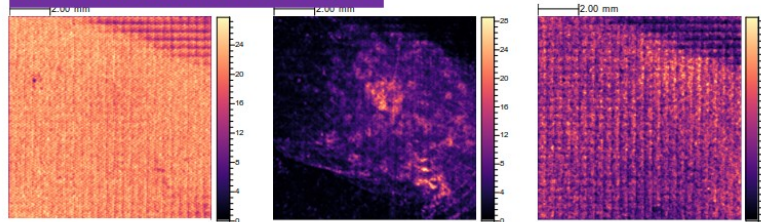


Cu^+

Cu_2OH^+

Cu^-

Surface contamination

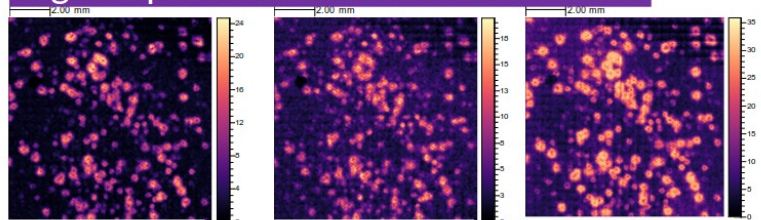


C_3H_5^+

Na^+

C_2^-

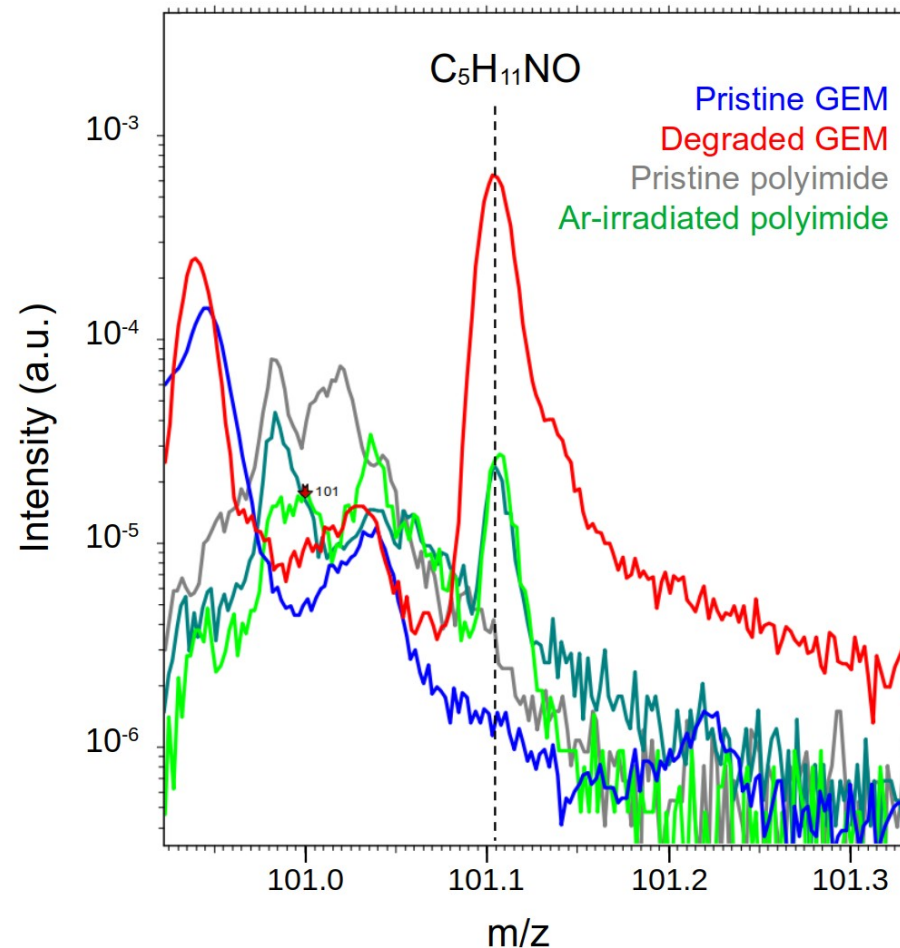
Organic pattern adsorbed onto surface



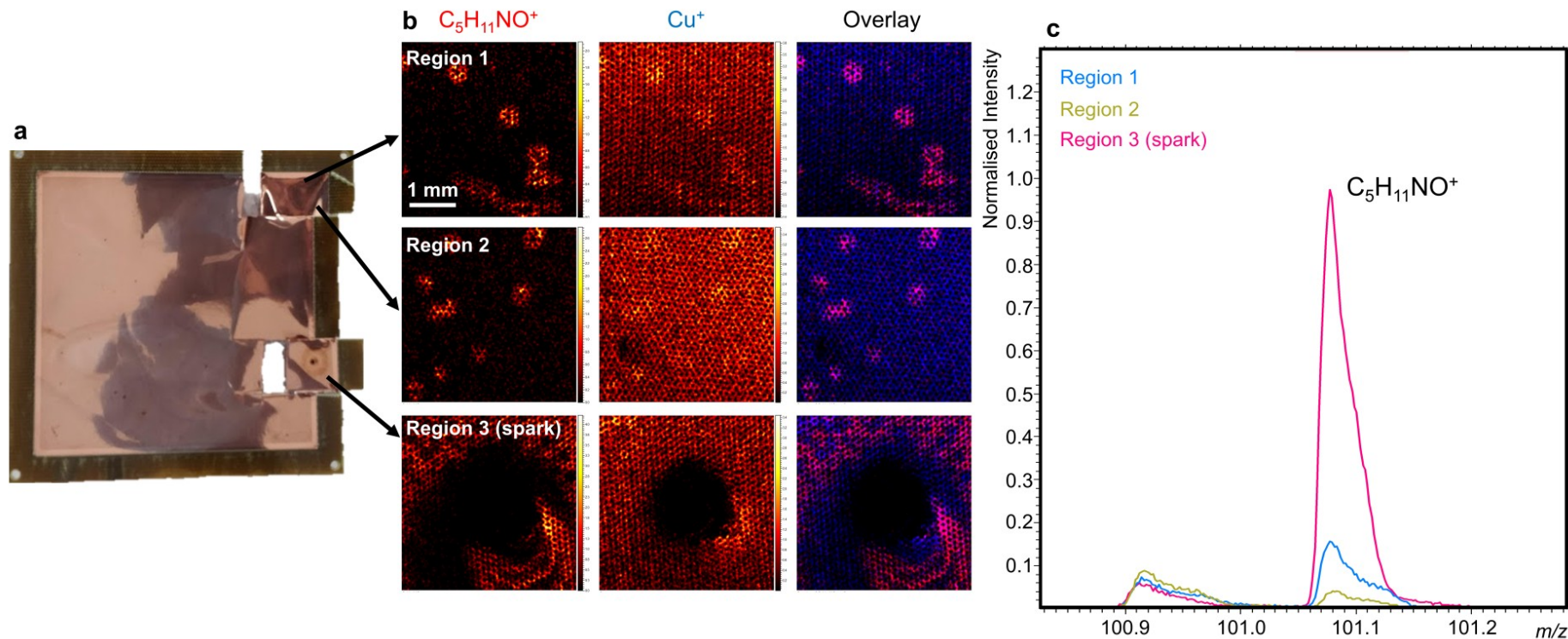
$\text{C}_5\text{H}_{11}\text{NO}^+$

CNCu_2^+

F^-

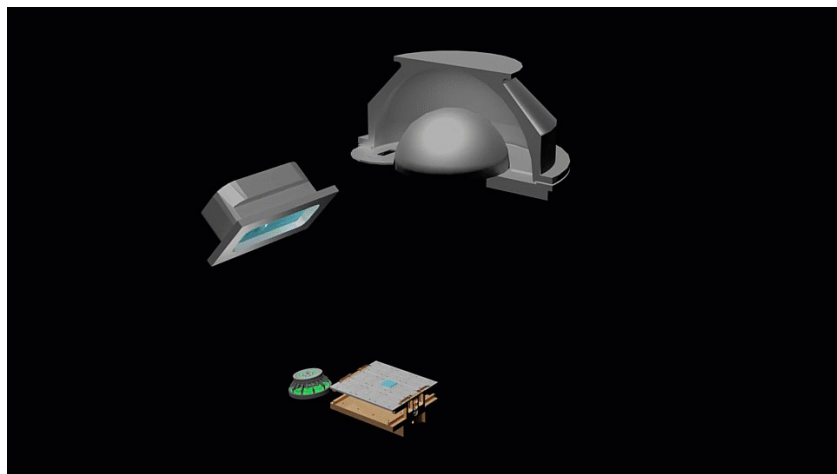


Possible connection with sparks!



X-ray Photoelectron Spectroscopy

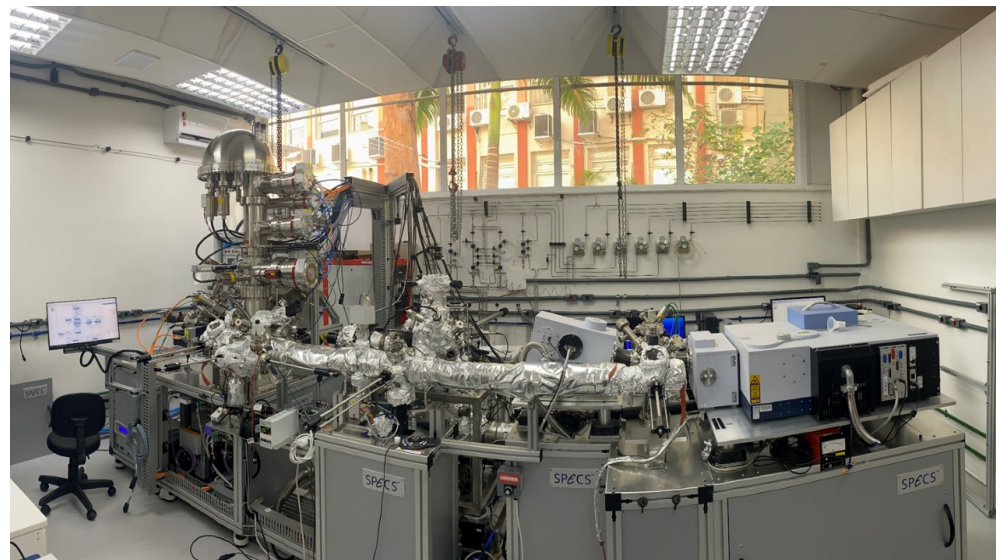
(XPS) works by irradiating a material with X-rays and measuring the kinetic energy of emitted electrons to determine the elemental composition and chemical states of the surface.



<https://ipgi.co.in/xps-x-ray-photoelectron-spectrometer/>

UHV system to NAP-XPS

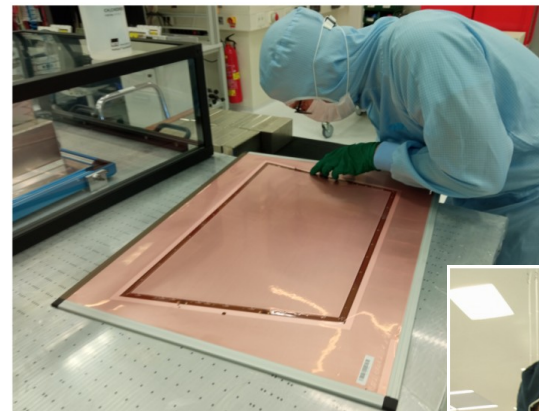
Lab. de Fenômenos de Interface e Superfície do CBPF



Near-Ambient Pressure X-ray Photoelectron Spectroscopy (NAP-XPS) analysis used to monitor the adsorption and surface chemical reactions between copper and CO₂ **(FOR DIFFERENT CO₂ PRESSURES)** in a GEM foil produced at CERN workshop.

- **GEM samples produced by the CERN workshop**
 - For the ALICE TPC spare parts
 - Stored in same conditions as the GEMs used in the assembly
- **Why not samples of copper?**
 - Adsorption is highly dependent of the surface conditions
 - Crystallinity, oxidation states, roughness, etc.
 - Using GEM samples as produced in the CERN workshop is meaningful as GEM samples

Sample preparation



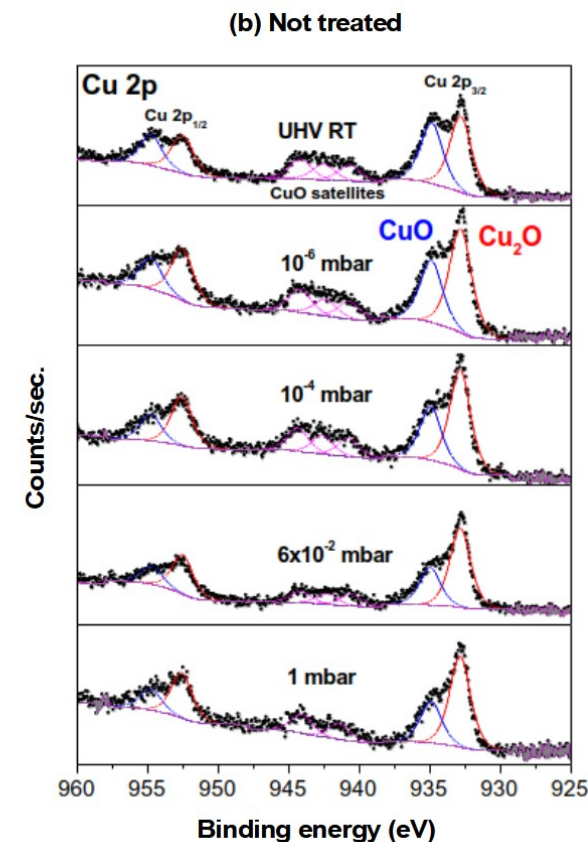
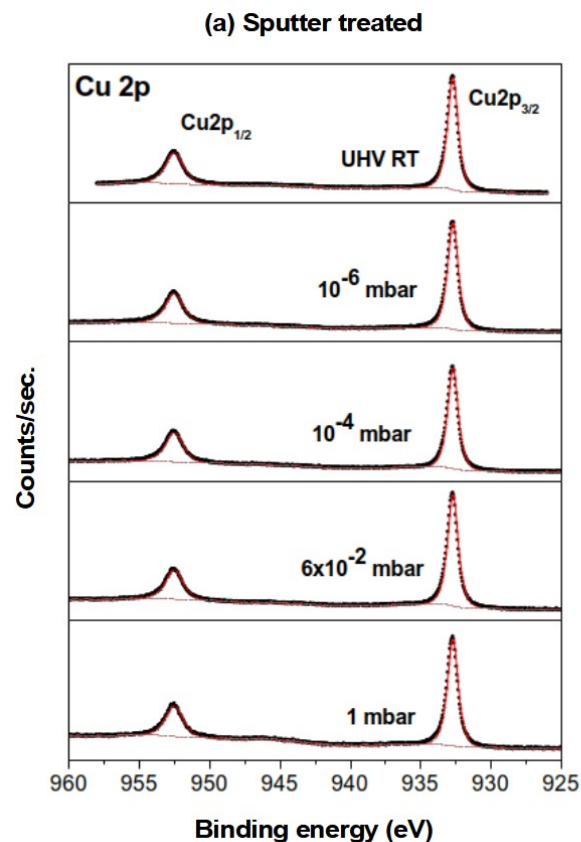
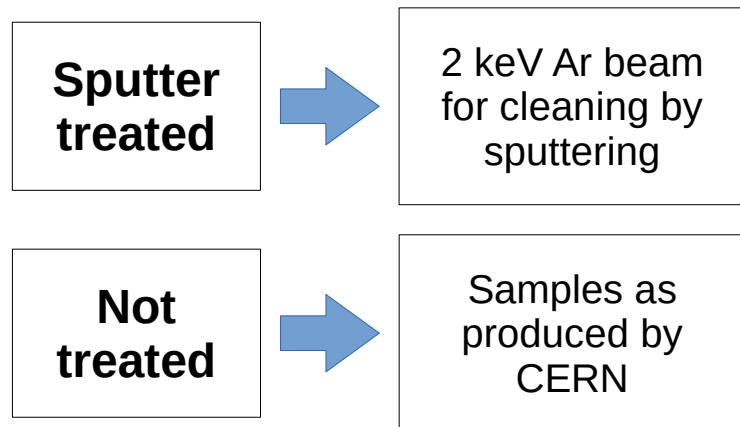
ALICE-TPC
samples from
spare mounts



T.B. Saramela
at the GSI lab

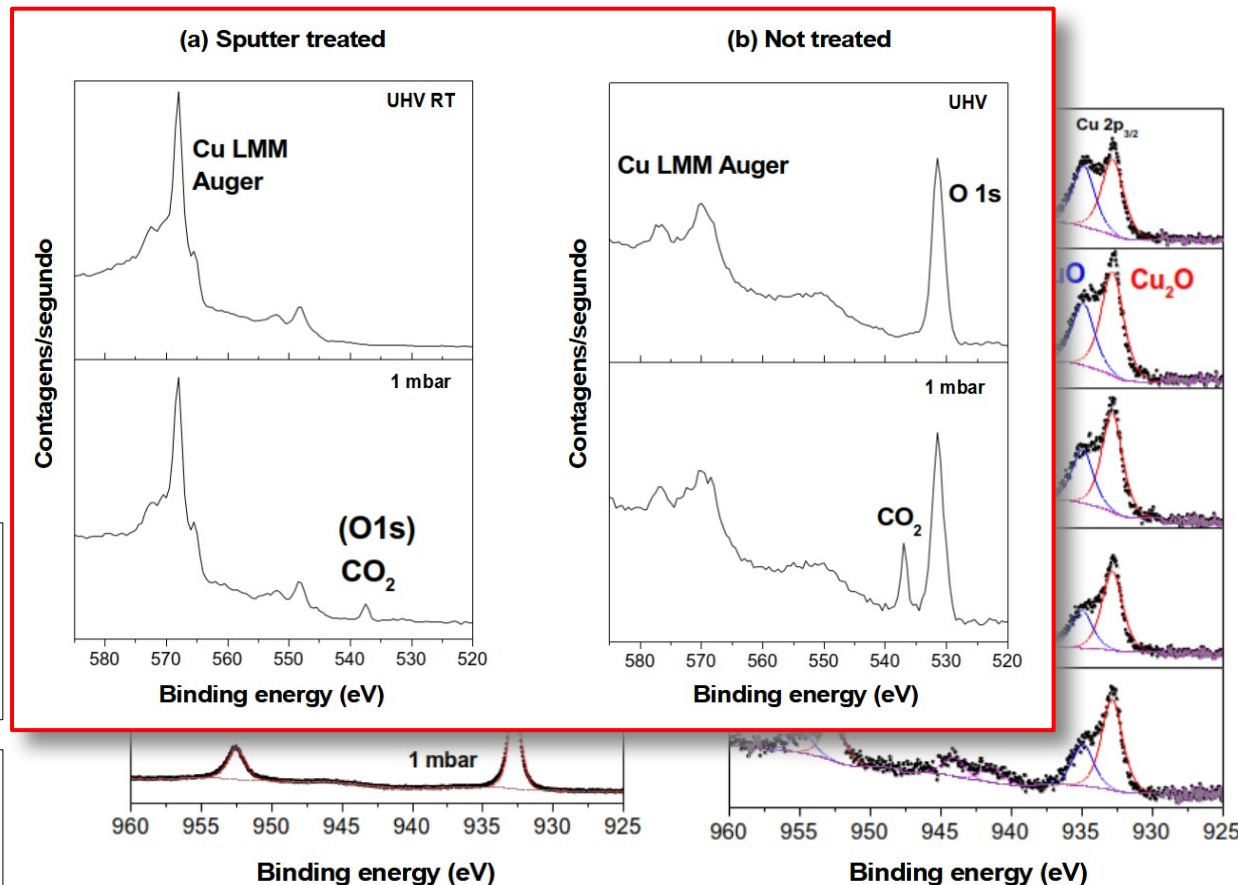
Results: the pristine surface is not metallic copper

- Copper signal from the surface for different CO₂ partial pressures
- **Pristine surface is a mixture of Cu₂O (Cu⁺¹) and CuO (Cu⁺²)**

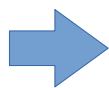


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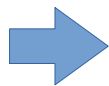


Sputter
treated



2 keV Ar beam
for cleaning by
sputtering

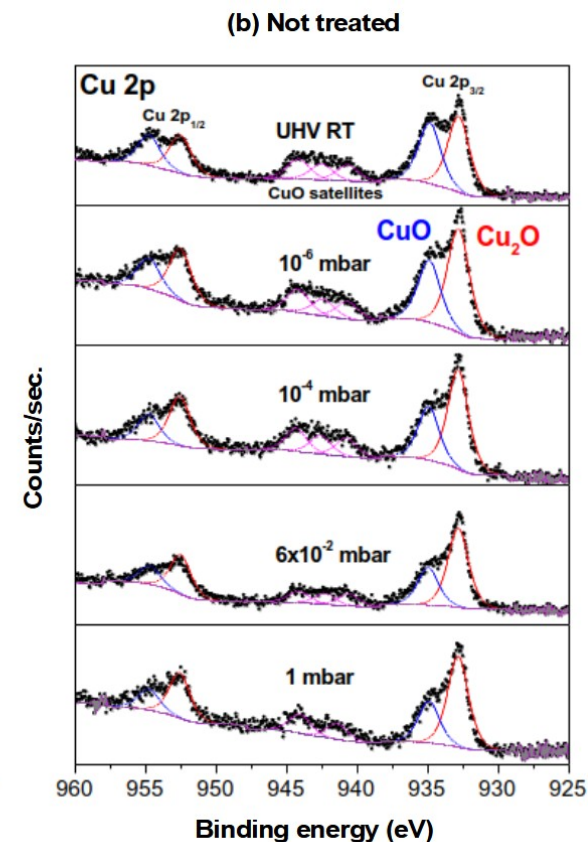
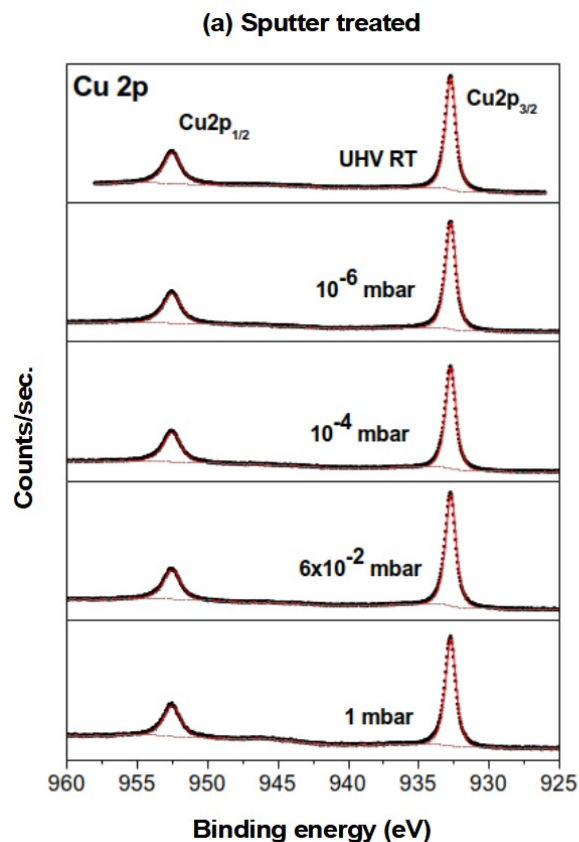
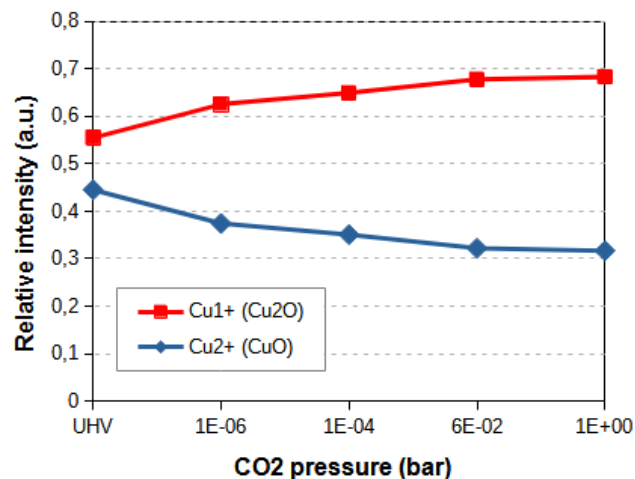
Not
treated



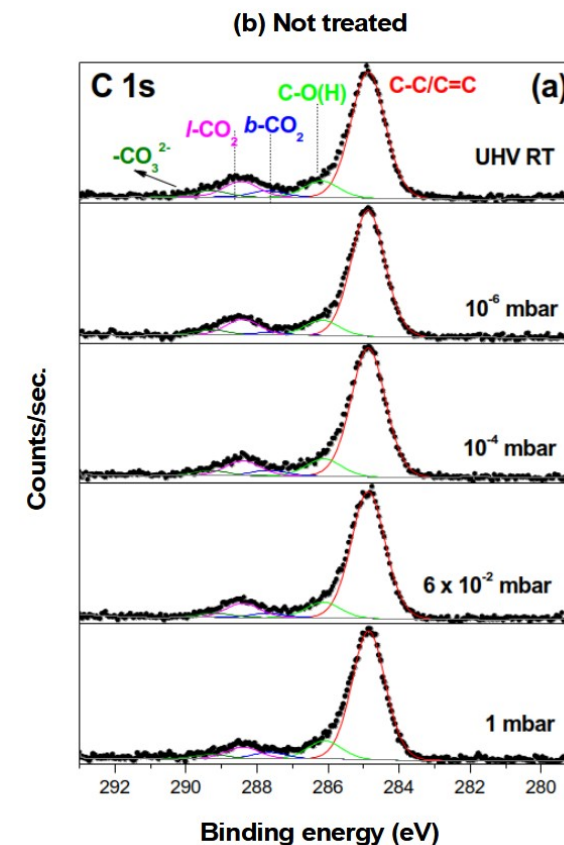
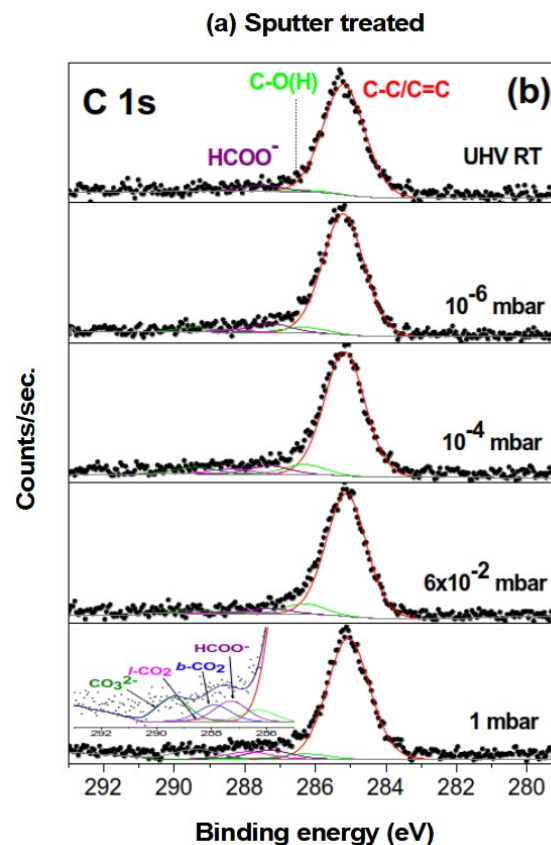
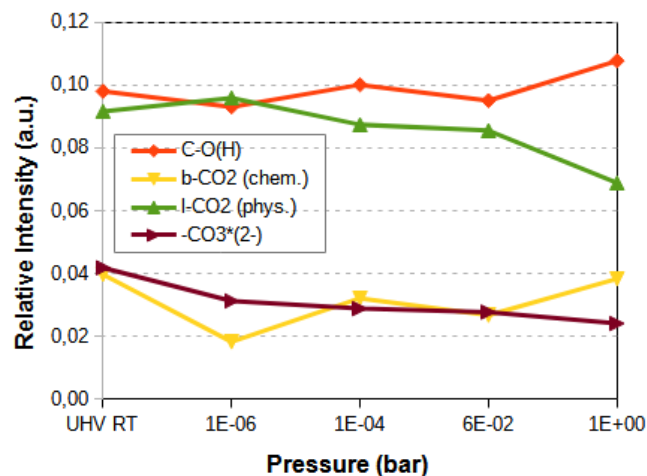
Samples as
produced by
CERN

Results: Cu_2O signal increases and saturates with pressure

- Copper 2p signal from the surface for different CO_2 partial pressures
- **CuO is converted into Cu_2O for increasing pressure, but saturates**

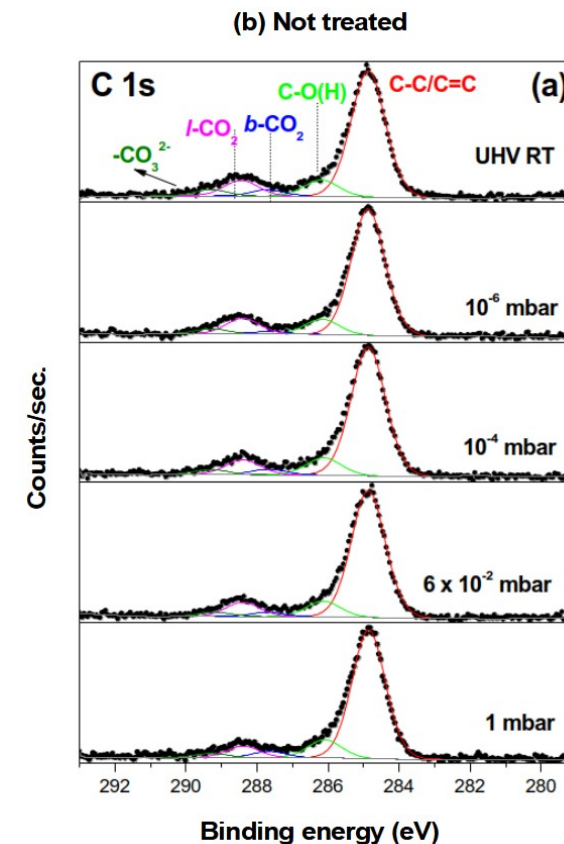
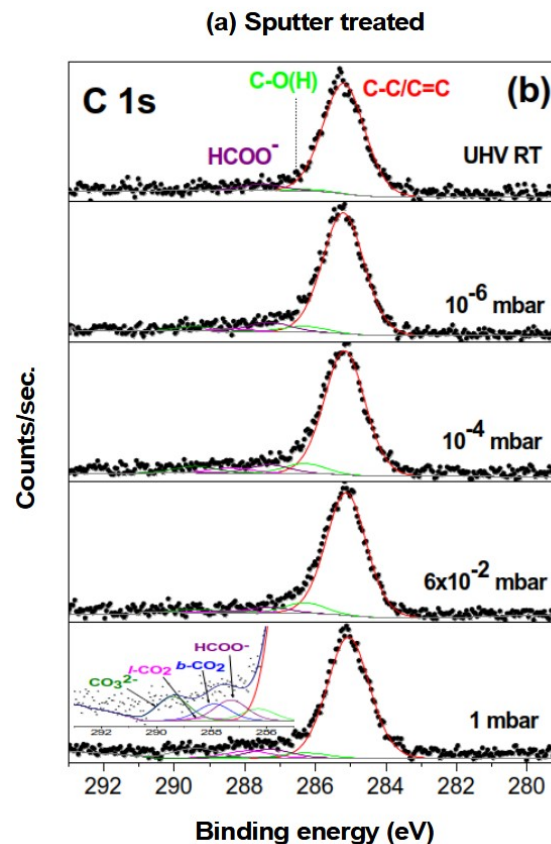
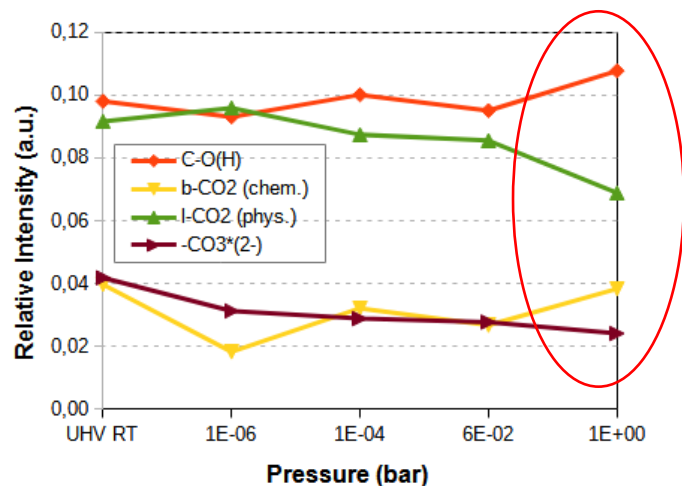


- Carbon 1s signal from the surface for different CO₂ partial pressures
- Carbonate, hydroxyl, physisorbed, and chemisorbed species form only on pristine samples.**



Results: Surface reactions on the pristine surface

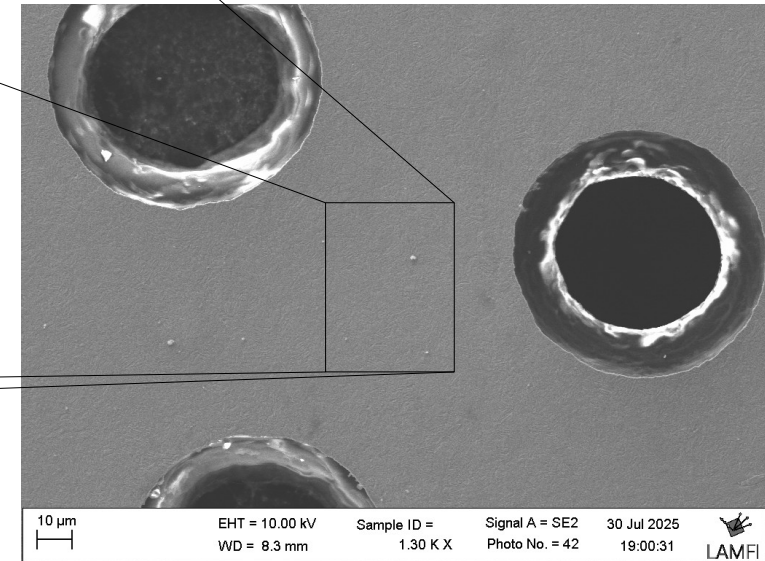
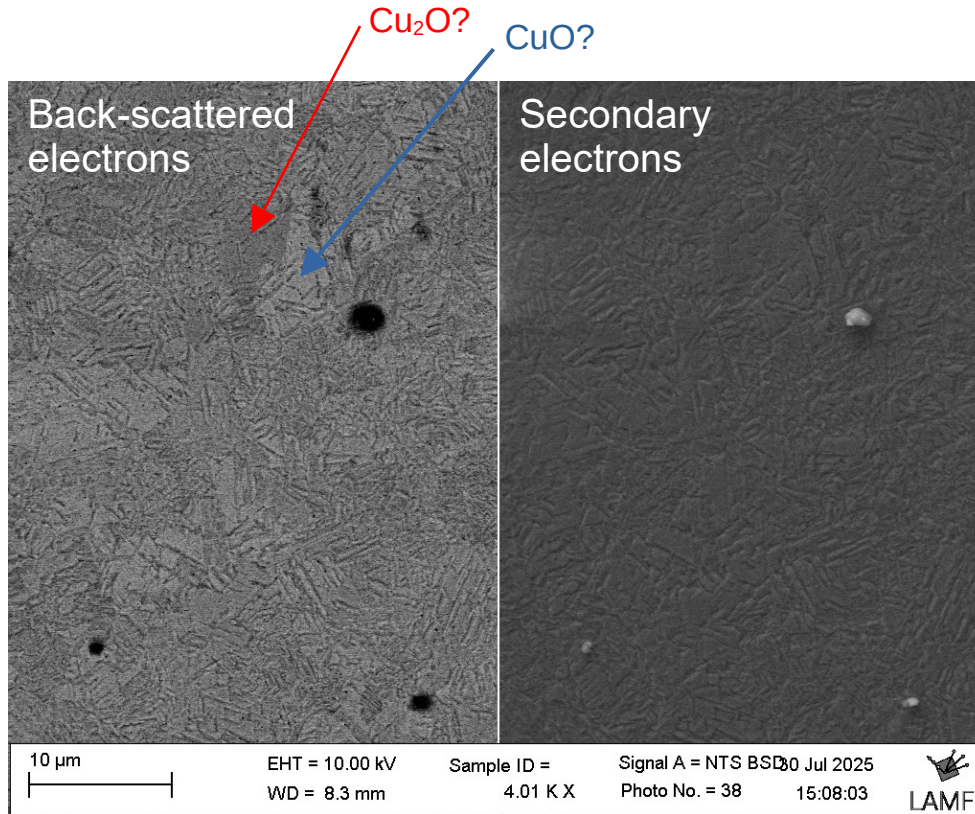
- Carbon 1s signal from the surface for different CO_2 partial pressures
- After Cu_2O saturation, l-CO_2 converts to b-CO_2 and the hydroxyl increases**



- **No metallic copper at pristine surface**
 - Metallic copper did not react with CO_2
- **CuO converts into Cu_2O**
 - CuO is not formed naturally (requires external energy)
 - May be **formed in chemical reactions** during production
- **Both, CuO and Cu_2O are p-type semiconductors**
 - 1.2 and 2.0 eV, respectively
 - **Not good insulators**, mainly in higher temperatures
 - Behave like conductors at 100 and 150 °C respectively
- **Cu_2O presents **low interaction** with CO_2**
 - Weak tendency to be an aging agent

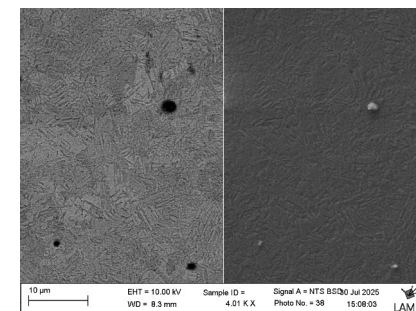
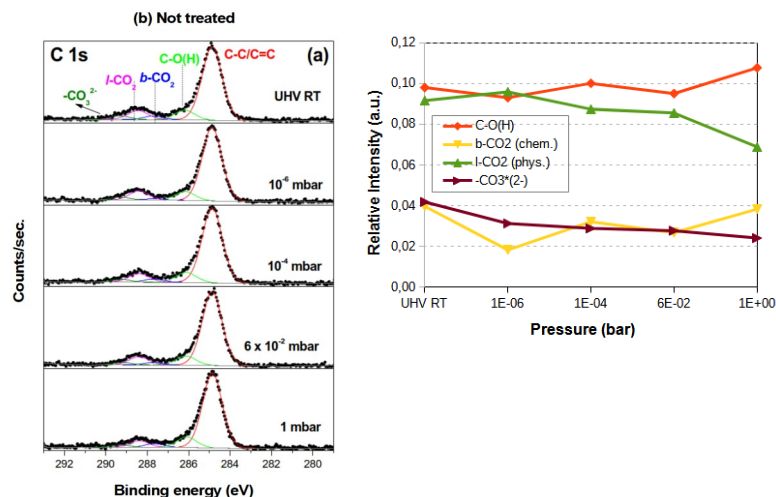
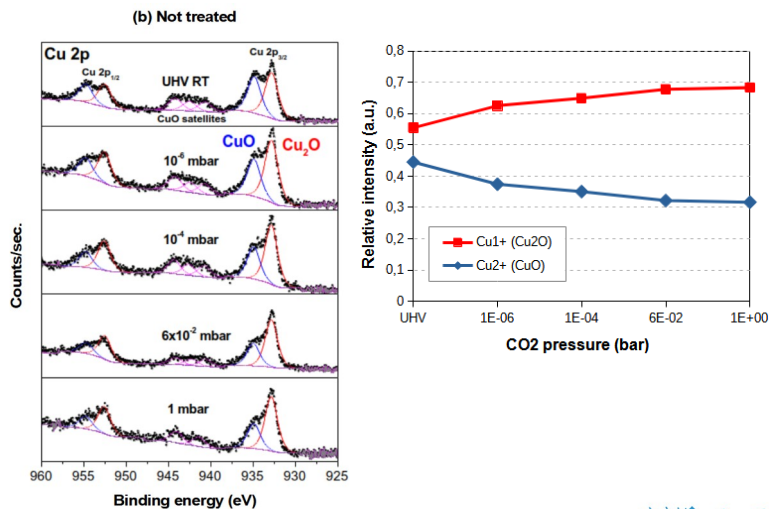
- Both pristine and sputter treated surfaces present carbon signals
 - No carbon reaction observed in sputter treated samples (graphite trace contamination?)
- Carbonate, hydroxyl, physisorbed, and chemisorbed species observed only on pristine samples
 - Carbonate signal reduces systematically in pristine sample
 - Hydroxyl presented increment after Cu_2O saturation
 - Dependent on adsorbed water (which is present in pristine samples)
 - I-CO₂ converts to b-CO₂ after Cu₂O saturation
 - Physisorbed CO₂ (Van-der-Waals bonding) become chemisorbed (covalent bonding)
 - Probably in the CuO residual region

Next steps: determine Cu oxidation states distribution



Unveiling Surface Chemistry and Ageing Pathways of GEMs through In-Situ Surface Spectroscopy

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