

Study of vacuum stability at cryogenic temperature

WP4 - Activity at LNF Alba 07-09/11/2017

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Beam Screen Temperature

Working Pressure \iff BS Temperature Range



Saturated vapour pressure from Honig and Hook (1960) (C2H6 Thibault et al.)

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- Find right working temperature is a fundamental point for vacuum stability.
- Work near a gas desorption temperature could generate great pressure oscillations.

Study of adsorption/desorption behaviour near critical temperature is mandatory to understand vacuum stability







Vacuum Stability









Studies

• Surface Quality

- X-Ray / UV Photoemission
- Secondary Electron Yield (SEY)

Adsorbed atoms and molecules

- Cryogenic temperature
- Atoms/Molecules Adsorption/Desorption process







LNF-Lab Now

Two Different Ultra-High Vacuum Systems equipped with:

- Low Energy Electron Diffraction
- Secondary Electron Yield Spectroscopy
- Surface Preparation
- Gas-Line

- X-Ray/UV Photoemission
- High Temperature Manipulator
- Low Temperature Manipulator (≈ 9 K)

Raman Spectroscopy

Scanning Tunneling Microscopy







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Gas-Line



HT Manipulator



LT Manipulator



LT STM





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LNF-Lab Activities I

Maintenance of Low Temperature System

Fixed some problems with Low Temperature manipulator, working on compressor of cryogenic system (≈5 months)

Now Low Temperature = 10 K







LNF-Lab Activities I





LNF-Lab Activities II

First **Preliminary** Results

Adsorption/Desorption process of Atoms and Molecules (Argon, CO) at cryogenic temperature measured with:

Secondary Electron Yield (SEY) and

Temperature Programmed Desorption (TPD)







- Adsorption process of Argon (Ar) and Carbon-Monoxide (CO) on atomically sputtered Cu surface at low temperature (with <u>SEY</u>)
- Desorption process of Ar and CO from heated Cu sample (with <u>SEY</u> and <u>TPD</u>)
- Interaction between electrons and Ar films (<u>SEY</u>)







LNF-Lab Activities II

Working Parameters:

- Basic Pressure $\leq 1 \times 10^{-10}$ mbar
- Electron Beam Current < 1x10⁻⁷ Ampere (A) (Max Current @ max electron energy)
- Electron Beam Energy from 75 to 1000 eV
- Sample Bias 75 V
- Single Spectra Acquisition Time ≈ 120 sec
- Beam Radius < 1.0 mm
- 1 Langmuir (L) = 1 sec @ 1x10⁻⁶ mbar
- 1 L = 1 Mono-Layer (ML) (with sticking coefficient = 1)
- Temperature range: form 10 to 300 K







• Why start with Atomically Sputtered clean Copper?

Electron Beams have not effect on a clean surface so any modification to the SEY can be attributed to atoms and molecules on surface.

- easy to single out contamination (sample pumping) from experimental.
- easy to eliminate spurious and otherwise occurring scrubbing effects from real non-clean surfaces.







LNF-lab Activities II

Experimental Test with Atomically Sputtered Copper



➢ First observable changes (blu) after 0.02C/mm² → more than 6 h SEY measurements

- > More significant changes (green) needs more than 60 h continuous measuring
- Our standard SEY lasts 120s!

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LNF-lab Activities II

- Why Argon and Carbon-Monoxide?
 - Argon is a inert gas and it's the best starting point to study SEY al Low Temperature
 - Carbon-Monoxide is a gas of great interest for accelerator physics







LNF-lab Activities II



Reference Literature spectra of Argon and Carbon-Monoxide SEY





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First Preliminary Experimental Results







Argon Adsorption (SEY)

uroCirCo



Argon Adsorption (SEY)







Argon Desorption (SEY)

Desorption process: Cu sample heated up to 100 K





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Desorption process heating Cu sample to 100 K





urcirCol



Argon Desorption (TPD)

Test of Temperature Programmed Desorption (TPD) with Mass Spectrometer



Three different peaks at 22, 30 and 37 K

due to the different desorption of TF from Manipulator (Peak 1 and TF/SL from sample





Argon Results

Adsorption Process (LT)

- Formation of a Thick Film (TF) at high coverage (increasing of SEY)
 - Formation of a Single Layer (SL) on Cu at LT (characteristic peaks in low energy region)

Desorption Process (Heating)

- System returns to the original state with slight differences
 - Possibility to follow formation of SL from TF
- Possibility to measure desorption temperature with SEY and TPD





Argon Adsorption II (SEY)



Second Run of Adsorption process of Argon on Cu sample at 10 K Same General behaviour







e⁻-Argon Interaction (SEY)

Sample Leaved under continuous scan as function of Time









e⁻-Argon (SEY)

SEY under continuous scan



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e⁻ bombardment induced different behaviour

- SEY at 930 decreases as a function of time
 - SEY at 10 eV remains constant





e⁻-Argon (SEY)



New Point presents different SEY spectra with the same features of SEY with large amount of Ar



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Argon Results II (SEY)

Beam-Layer Interaction

Argon Thick film interacts with primary electrons and *desorbs*









CO Adsorption (SEY)

Adsorption process of Carbon Monoxide on Cu sample at 10K General behaviour







CO Adsorption (SEY)

Adsorption process of Carbon Monoxide on Cu sample at 10K Low Energy behaviour





CO Adsorption II (SEY)

Adsorption process of Carbon Monoxide on Cu sample at 10K General behaviour



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- Higher total Langmuir
 exposure
- Same characteristic features of TF and SL







CO Desorption (SEY)

Desorption process of Carbon Monoxide on Cu sample heated up to 300 K General behaviour



- Not Linear behaviour as in Ar
- Two different
 Desorption process
 around 30 K and 200 K
 respectively







CO Results (SEY)

Adsorption Process (10 K)

• Formation of a TF with Low SEY

Characteristic peaks for SL in the Low Energy Regions







Work Function (SEY)

Work Function variation



No changes in the first steps of adsorption

Changes after formation of the multi-layer



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- Follow Adsorption process checking the High-Energy SEY (HE-SEY)
- Distinguish Single Layer (SL) formations from Thick Film (TF) by Low-Energy SEY (LE-SEY) behaviour
- Quantify the numbers of adsorbed gas layers on surface
- Measure the desorption temperature of TF and SL with SEY and TPD
- Measure Work Function (WF) variation







Future Activities

- Different Gases adsorption (CO, CO2, CH4 ...) (pure and mixture)
- Electron desorption
- Thermal Programmed Desorption (TPD)

- Gas-Line Upgrades
- TPD Upgrades







Upgrade Gas-line on low temperature system
 Low pressure gas dosing in front of sample (with gas line upgrades)

Upgrade TPD system

Measure of temperature, Heating system, relative position in front of mass spectrometer



