

## Vacuum Technology for Synchrotron Radiation Sources

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#### About Vacuum

#### Vacuum = Empty

AVS definition: "Vacuum refers to a given space filled with gas at pressures below atmospheric. Molecular density less than about 2.5x10<sup>19</sup> molecules/cm<sup>3</sup>

#### P = nkT [mbar)

n ... number of molecules per unit volumek ... Boltzman's constantT ... temperature in Kelvin



#### I About Vacuum

#### ☐ Vacuum ranges

Low (and medium): from atmospheric to about 10<sup>-2</sup> mbar Piston pumps, water ring, rotary, sorption (nitrogen)

High vacuum: from 10<sup>-3</sup> mbar to 10<sup>-7</sup> mbar Roots, ejector, diffusion, molecular (water vapour)

Ultra-high vacuum: from 10<sup>-8</sup> mbar to 10<sup>-16</sup> mbar Ion, cryogenic, sorption (hydrogen)



## I About Vacuum Pumping

Main parameters:
→ the lowest pressure (ultimate pressure)
→ the pressure range
→ the pumping speed
↓ S = V/t [1/s]
↓ S is pumping speed, V is the volume, t is time
→ the exhaust pressure (additional pump)
→ the selectivity and residual gas composition (UHV)



## I About Vacuum Conductance

**Conductance** C [l/s]  $N = C(n_1 - n_2)$ N ... number of molecules  $n_1, n_2$  ... concentrations on both part of the conductance C depends on the shape and geometry of the components and on the type of gas (analytical or numerical calculations)



I About Vacuum Outgassing

Desorption rate from materials -Arrhenius' equation dN/dT = - const. N [exp(-E/kT)]\_ E ... binding energy of the molecules on the surface [kJ/mol] dN/dT = Q ... thermal outgassing rate \_ At the equilibrium (UHV systems)  $\square P = Q/S$ 



#### II Vacuum requirements for synchrotron radiation sources

Beam lifetime due to gas density must be > 10 h  $1/\tau = 1/\tau_{el} + 1/\tau_{br} + 1/\tau_{tousch}$ Short "conditioning time" **Quick recovery after venting** Simplicity in modification for new installations Smooth chamber wall design **ELETTRA** - operating pressure < 10<sup>-8</sup> mbar (dynamic pressure < 10<sup>-10</sup> mbar/mA)



#### RF shielding - "fingers"





## III The choice of material

\_ Sufficient mechanical strength \_  $L_c = 1.11 \text{ D(D/h)}^{1/2}$ 

⊥ L<sub>c</sub> ... critical length for cylindrical parts
 ⊥ D ... mean diameter
 ⊥ h ... wall thickness

**\_ Impermeable enough to gases** 

∟ Low vapour pressure (Cd, Zn - no!!!)



## III The choice of material

**Good resistance to special working** conditions (temperature, humidity) Low specific outgassing rate **Low desorption yield \_** Low magnetic permeability Good weldability and machining Good thermal conductivity **Required electrical parameters** 



## The choice of material

#### The mostly used materials:

- J stainless steel (bad thermal conductivity)
- □ aluminum (porous material, high desorption yield coef.)
- $\Box$  copper (soft)
- 」 titanium véry promising material (expensive)

□ Coated chambers( Cu, TiN, NEG - Non Evaporable Getter)

Getter: a material which is included in a vacuum device
 (tube) for removing gas by sorption

NEG: Ti-Zr-V alloys, activation at 150-180 °C for 24 h do not pump methane and inert gases!



### IV

## **Cleaning procedures**

Necessity to remove impurities and hydrocarbons
 Specific outgassing rate:

 $\Box q_D < 1.5 \ge 10^{-12} \text{ mbar l/s cm}^2$ 

prior to assembly after assembly

- → physical cleaning (abrasives)
- ☐ chemical cleaning (solvents)
- ☐ firing at ambient or inert (???)

plasma cleaning (O<sub>3</sub>, N<sub>2</sub>) glow discharge cleaning in-situ bake-out atmosphere



## Vacuum Chamber Design

J Main parts of the ring vacuum chamber:
 J - electron chamber (elliptical, cylindrical, rhomboidal)

- bending magnet chamber: antechamber solution
   efficient removal of desorbed gases far away from the
   electron trajectories through the slots
- ☐ with or without central pumping
- ☐ (NEG coated chambers lower conditioning time)



#### **Pumping spot for the electron rhomboidal chamber**





#### **Aluminum Bending Magnet Vacuum Chamber**





#### Insertion device vacuum chamber with central pumping





#### \_ UHV conditions:

 $\square$  - static pressure (without beam) < 10<sup>-10</sup> mbar

J - operating pressure (with beam) < 10<sup>-8</sup> mbar
 J Total gas load

 $Q_t = Q_{des} + Q_{ind}$ The total photon flux (Souchet at all, 1983)

 $\ \ \, \square \ \, dN_{ph}/dt = 8.08 \ \, x \ \, 10^{17} \ \, I \ \, E \ \, [photons/s]$ 

- □ I ... beam current in mA
- □ E ...machine energy in GeV

# elettra

## VI Pumping Requirements

ELETTRA: 400 mA at 2 GeV,  $\rho = 5.5$ m Total number of photons =  $6.4 \times 10^{20}$  ph/s ▲ Molecules extracted from the walls  $\Box N_{mol} = \eta N_{ph}$  $\square \eta \dots$ desorption yield coefficient Characteristic of materials - has to be measured The  $\eta$  decrease causes the pressure decrease, so called "conditioning" Measured against Amperhour (integrated current)





Pumps must be uniformly distributed along the vacuum chamber ☐ Pressure profile calculated at steady state  $dO = C L d^2P/dx^2 dx$ <u>L... half distance between two pumps</u> □ C ... pipe conductance of the perimeter B Maximum pressure  $\square P_{L} = q_{D} B L [1/S_{P} + 1/2C]$ Minimum pressure  $\square P_0 = q_D B L / S_P$ 



# elettra

## VI Pumping Requirements

☐ The equilibrium pressure in the ring
 ☐ P = (17.8 η + q<sub>D</sub> A) S<sub>ef</sub>
 ☐ A ... the internal surface of the volume V
 ☐ Compromise between C and A (experience)



#### **Cross Section of the Extruded Aluminium Insertion Device Vacuum Chamber with Cooling Chanels**





#### Aluminium Insertion Device Vacuum Chamber without central pumping





## VII Beam cleaning efficiency

□ The third generation synchrotron light sources
 □ 1 Ah of conditioning = 1.7x10<sup>23</sup> ph/m (R.P.Walker)
 □ η in the range of 10<sup>-7</sup> mol/ph can be reached after about 5 Ah of conditioning
 □ After about 10 Ah of conditioning the specific outgassing rate is < 1x10<sup>-13</sup> mbar l/s cm<sup>2</sup>
 □ Beam is the best cleaning agent !



## VII Beam Cleaning Efficiency

In situ bake-out - described as flash desorption  $\Box E_{\rm D}/RT_{\rm m}^2 = \text{const. exp}[-E_{\rm D}/RT_{\rm m}]$  $\Box$  E<sub>D</sub>... activation energy of desorption  $\Box$  T<sub>m</sub>... surface temperature at the maximum desorption rate  $\Box E_{D} = 300 \text{ x } T_{m} \text{ [kJ/mol]}$ Energy of photons  $E = h v = h c/\lambda$ ☐ h, c ... Planck's constant, velocity of light, resp.  $\neg$   $\nu$ ,  $\lambda$  ... frequency and vawelength of photons



## VII Beam cleaning efficiency

□ Photon "cleaning"□ λ [nm] 500 200 1□ E [eV/mol] 4x10<sup>26</sup> 8x10<sup>26</sup> 2x10<sup>27</sup>



## VII Beam cleaning efficiency

#### **LETTRA** experience:

- $\perp$  1) no complete bake-out before start up (1993)
- ☐ 2) enormous thermal stress leaks found
- □ 3) residual mass spectra: 2(H<sub>2</sub>), 16(CH<sub>4</sub>), 18(H<sub>2</sub>O), □ 28(CO, N<sub>2</sub>), 44(CO<sub>2</sub>)
- → 4) no difference in conditioning time between baked
- and unbaked vacuum sectors
- ☐ Bake-out is time consuming, non negligible cost



## VIII Pumping system

 Sputter ion pump: used crossed electric and magnetic field
 Consists of: stainless steel vessel anode of honeycomb construction titanium cathode magnets

Advantages

Disadvantages

J Gas captured inside
 J High S for active gases
 J Clean vacuum
 J No vibrations

Start to work at 10<sup>-5</sup> mbar Low S for inert gases (He) Argon instability (triode) Need refreshing (at 220 °C)

#### Pressure measurements according to the SIP's current reading



Current absorbed by the pump I = K P<sup>n</sup>

**Calibration equations** 

$I = 1850 P^{1.65}$	<b>SIP 45 l/s</b>
$I = 1740 P^{1.18}$	<b>SIP 60 l/s</b>
$I = 1590 P^{1.06}$	SIP 120 l/s
$I = 1260 P^{1.1}$	<b>SIP 230 l/s</b>
$I = 1200 P^{0.99}$	<b>SIP 400 l/s</b>
$I = 1050 P^{1.03}$	SIP 900 l/s



## VIII Pumping system

Ti sublimation pump
 Main parts: water (or LN<sub>2</sub>) cooled SST vessel
 wire feed spool driven from outside
 Ti wire
 heated post
 filaments (cathode)
 Do not pump helium!



#### IX Pressure measurements



Fig. 4.64 Range of pressures covered by vacuum gauges.



IX Pressure measurements

Ionisation gauges - "hot cathode"

#### Bayard-Alpert hot cathode gauge



I<sup>+</sup> = I <sup>-</sup> s P from 10<sup>-3</sup> to 10<sup>-11</sup> mbar Need calibration for each gas species Linear calibration curve Sensitivity variation Outgassing



#### VIII Pressure measurements

Ionisation gauges - "cold" cathode

#### Some "Spurious" Effects



Crossed electric and magnetic field Self sustaining magnetic discharge Wide range - IMG Inverted magnetron gauge 10<sup>-3</sup> - 10<sup>-11</sup> mbar

Need calibration for each species Striking at low pressures





#### Residual gas analyzer



Applied d.c.+a.c. for given frequency e/m is detected

Leak detector only mass 4 (He) registered



#### IX Residual Gas Analyses

**OMA characteristics:** 1) high pressure performance (< 1x10<sup>-5</sup> mbar) 2) detection of small signals - fragment ions detection 3) dependence of sensitivity on gas species 4) hysteresis - less pronounced at  $P > 10^{-8}$  mbar 5) formation of fragment ion 6) degassing of vacuum heads 7) total pressure measurement 8) last but not least - COSTS



## X Reliability under real experimental conditions

#### What is special:

- very big facility
- must operate reliably
- radiation
- electric field
- external magnetic field
- RF structures
- overheated parts of the vacuum chamber

There is no space for vacuum instrument installation!



## X Reliability under real experimental conditions

How to cure effect of:

 radiation: developed a metal structure installed with the PEG
 magnetic field: special shielding for PEG μ metal saturates over 300 Gauss double ARMCO sheets for RGA
 RF structure: inside the RF cavities frequency shifter installed



#### Detail of the pump attachment on the bending magnet vacuum chamber





#### Radio Frequency Cavity from Cu with cooling channels





## Bending magnet with the vacuum chamber





#### Photon absorber







# Design of the accelerator vacuum system is not only the science

It is an art