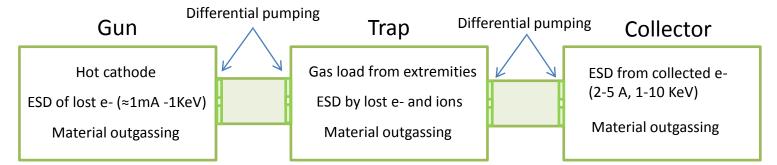
HIE-EBIS Workshop Vacuum Aspects and Recommendations

Paolo Chiggiato CERN Technology department Vacuum, Surfaces and Coatings group Injectors & Vacuum Measurements

Request to vacuum system



Pressure requirements							
 Protection of cathode Reduced gas transmission 	- Lowest possible gas density	- Reduced gas transmission					
10 ⁻⁹ mbar	$10^{\text{-}13}mbar$ Is this value mandatory also for $\mathrm{H_2}?$	10 ⁻⁸ mbar					

Gas load in the trap from collector chamber without differential pumping:

$$Q = P_C C_{C \to T} \sim 10^{-8} \text{ mbar } \frac{\ell}{s} (C_{C \to T} \sim 1 \frac{\ell}{s} \text{ for cylindrical tube D=1 cm, L=10 cm})$$

Necessary pumping speed in the trap:

$$S_T = \frac{Q_{C \to T}}{P_T} \sim \frac{10^{-8}}{10^{-13}} = 10^5 \frac{\ell}{s} \to \text{irrealistic}$$

→Differential pumping is necessary

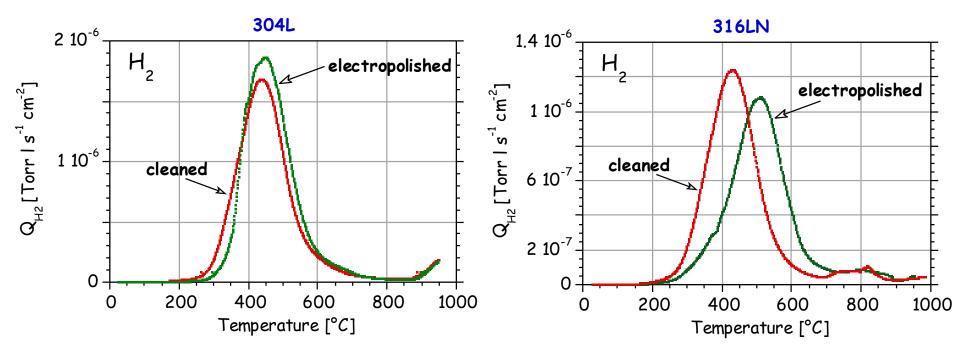
16 October 2012

P. Chiggiato CERN HIE-EBIS workshop

Gas sources: outgassing

Thermal outgassing for traditional vacuum materials after bakeout:

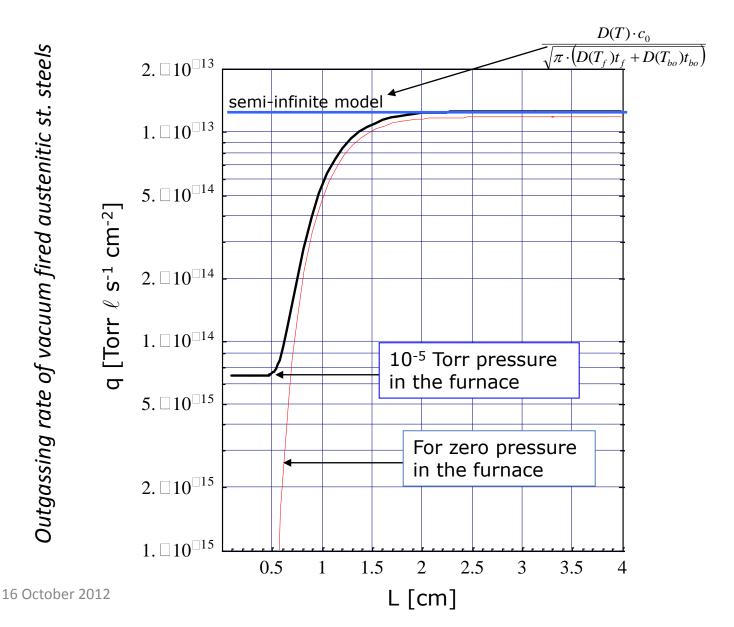
- Mainly H₂, except for porous solids (ceramics and graphite)
- Low T (<200°C) bakeout: 10⁻¹² mbarls⁻¹cm⁻² for stainless steel and copper, lower for Al alloys
- **Chemical and electrochemical** treatments have **minor effects** on H₂ outgassing rate.



Thermal desorption measurement: 1 mm thick ribbons, heating rate 5 K min⁻¹, bakeout temperature 150°C (15h)

 $P = \frac{\mathbf{v}}{\mathbf{s}} + P_0$

- Stainless steel vacuum fired at 950°C (2h): 2 10⁻¹³-7 10⁻¹⁵ mbar ls⁻¹cm⁻²



Effect of temperature on outgassing rate

$$Q \propto e^{-\frac{E_D}{k_B T}}$$

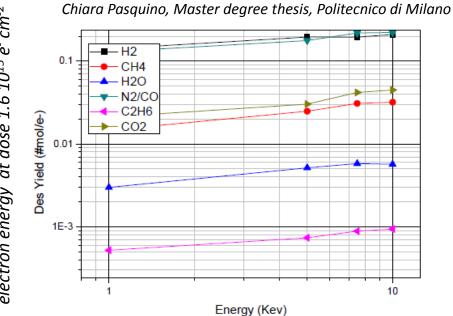
Example: outgassing rate of stainless steel

$$\frac{Q_T}{Q_{RT}} = e^{-\frac{0.52}{k_B} \left(\frac{1}{T} - \frac{1}{T_{RT}}\right)}$$

T = 100 K →
$$\frac{Q_T}{Q_{RT}}$$
 = 4 10⁻¹⁸
T = 300°C → $\frac{Q_T}{Q_{RT}}$ = 21000

Gas sources: ESD





 $P = \frac{\mathbf{Q}}{S} + P_0$

In general: the electron energy has a minor effect on η for electron energies higher than 2 KeV.

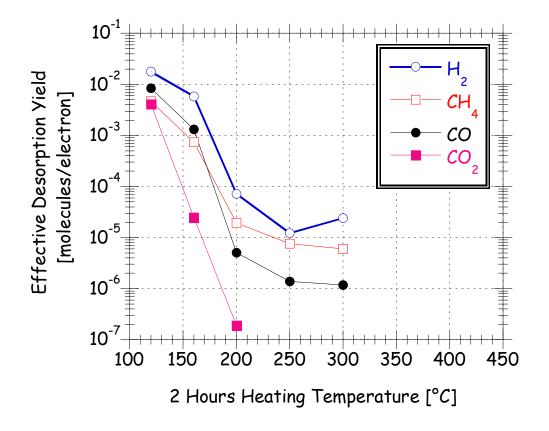
$$\eta = \eta_0 D^{-\alpha} \qquad 0.6 < \alpha < 1$$

Example:

- η=10⁻¹ at 10¹⁷ e⁻ cm⁻²
- α =0.6 and electron current 3.5 A
- Electron energy \sim KeV

Conditioning time	1 day	10 days
Desorption yields	1.6 10 ⁻⁵	4 10 ⁻⁶

- The desorption yield is only slightly affected by chemical and thermal treatments.
- Only TiZrV thin film coating may provide low effective desorption yields without conditioning



Gas pumping

$$P = \frac{Q}{S} + P_0$$

Pumps with P_o lower than 10⁻¹² mbar are needed

Getter pumps \rightarrow hydrogen dissociation pressure.

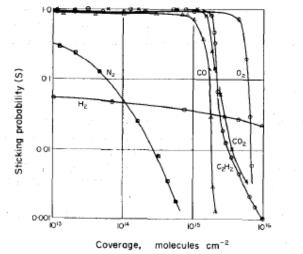
Hydrogen is pumped reversibly. In an isolated system an equilibrium pressure is obtained. It is never a problem in the UHV range.

Cryopumps \rightarrow saturated vapour pressure Vapour pressure below 10⁻¹² mbar for all gases, except He, may be produced at a temperature of 2.3 K, which may be obtained by means of a liquid He bath at reduced pressure (about 50Torr).

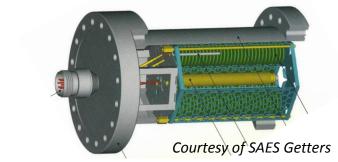
Lump getter pumps

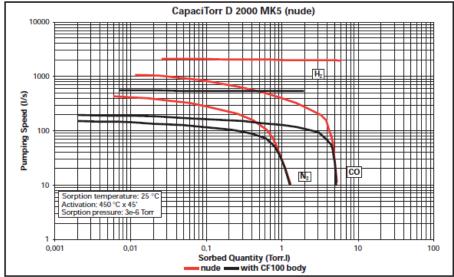


A K Gupta, J H Leck, Vacuum, 1975



- Saturated after the adsorption of one monolayer (3.3x10⁻⁵ Torr I cm⁻²) of CO.
- Already tested at 10⁻¹³ mbar

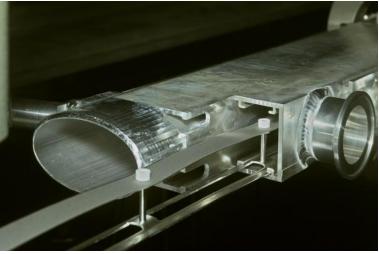




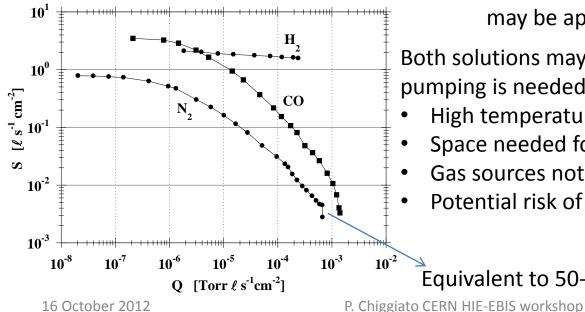
- Q_{sat}= 5 Torr ℓ equivalent to 15 m² of smooth getter surface.
- Already tested at 10⁻¹² mbar

Getters do not pump rare gases and CH₄: they always need auxiliary pumps

Getter pumps: NEG ribbon



St 101 In LEP, it was activated by ohmic effect





St 707

This alloy may be activated at temperatures as low as 300°C (24h): passive activation may be applied

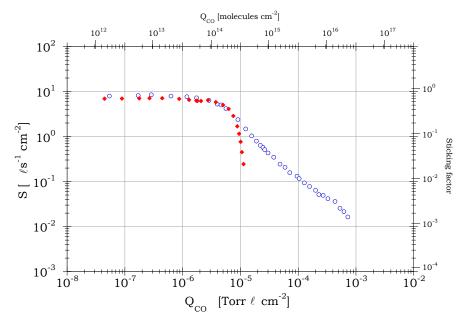
Both solutions may be used when distributed pumping is needed, however:

- High temperature needed for activation (T>300°C)
- Space needed for supports
- Gas sources not reduced
- Potential risk of dust production

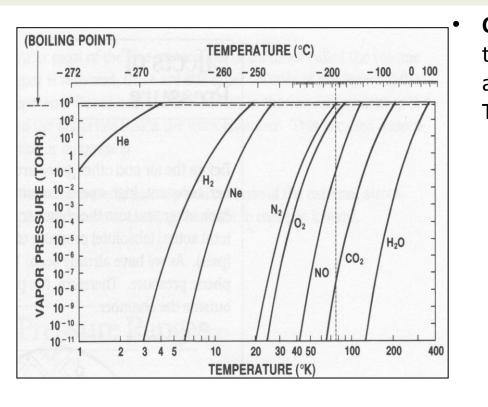
Equivalent to 50-100 cm² of smooth getter surface

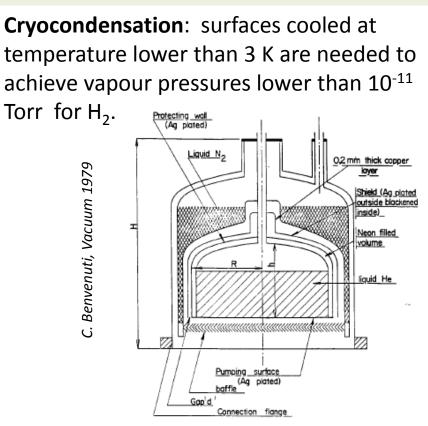
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- TiZrV is deposited by sputtering onto the inner wall of vacuum chambers.
- It may be activated at 180°C (24h). It is the main pump of the warm LHC's LSS.
- Typical S at zero coverage: 0.3-0.5 ℓ s⁻¹ for H₂.
- The pumping capacity may be increased by coating rough coatings on rough substrates
- Very large surfaces may be coated
- The outgassing rate of the substrate surfaces is suppressed
- No experience in pumping at cryogenic temperature, but we are eager to start...



Cryo pumps





- **Cryosorption:** submonolayer pumping → pressures much lower than the vapour pressure are achievable.
- Porous materials are used; for example, in one gram of standard charcoal for cryogenic application, about 1000 m² of surface are available for adsorption.
- Significant quantities of H₂ and He may be pumped at 20 K and 4.3K, respectively.

Cryo vs NEG pumps

	NEG pumps	Cryosorption pumps			
Pumping selectivity	No pumping for rare gas and CH ₄ ; need of auxiliary pumps	No selectivity			
Regeneration/Condi -tioning	Full bakeout for passive activation of NEG films or St707 ribbon	Auxiliary pumps to remove desorbed gas.			
Maintenance	No maintenance	Regular and expensive for refrigerators			
Power cut	Continue pumping, no need of power supply	Desorption; need of an auxiliary pump.			
Bakeout issue	Temperature of NEG surfaces > 180°C	Need for bakeable cryosorbers			
Cost	Lower	Higher			
Dust production	Not excluded a-priori for ribbons	No problem			
Safety issues	Pyrophoricity, except for NEG films	No problem for refrigerators			
Space needed	Negligible	Integration of refrigerator			
Inlet valve for lump pumps	No need	Necessary for regeneration -> €			
Operational incidents	Strongly affected by contamination (F, Cl, hydrocarbon)	Unduly gas charge on porous surface			

- Nowadays, no commercial vacuum gauges can measure 10⁻¹³ mbar at room temperature
- At CERN, a Bayard-Alpert gauge developed in the '70th is currently used.
- It has a x-ray residual pressure of about 2 10⁻¹² Torr; pressure measurement in the 10⁻¹³ Torr range is possible by modulation.
- The modified Helmer gauge gave reliable results in the low 10⁻¹⁴ Torr: but it is not anymore available at CERN. A development programme is going to be launched in 2013 in collaboration with an industrial partner.

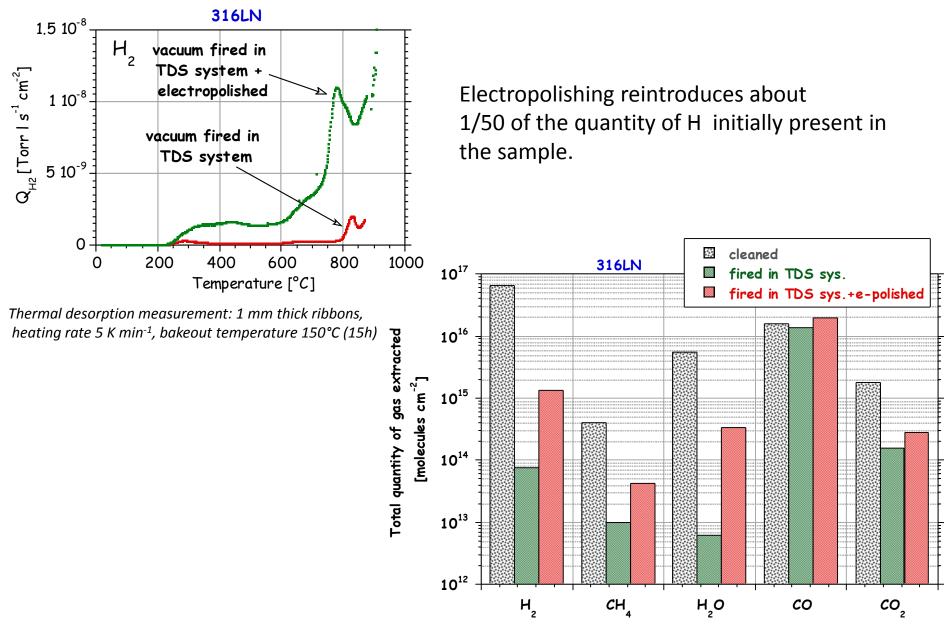


Conclusions

- The pressure requirement in the HIE-EBIS's trap is so challenging that every **single** material must be selected, treated and checked with the highest precaution.
- If a getter based solution is selected, be sure that:
 - the coated surface is the largest possible;
 - the design allow a uniform temperature bakeout;
 - during the activation, the coated surface is not unduly charged by gas desorbed by uncoated areas;
 - no F contamination is present (for example Teflon in primary pumps)
 - the NEG is activated after the full conditioning of the collector and cathode.
- If a cryosorption solution is selected, be sure that:
 - all parts are bakeable;
 - the cryosorber is strongly fixed and can withstand several bakeout cycles;
 - consider a scenario that include power cut;
 - cool the cryosorber surface as late as possible after the beginning of the pumpdown.
- As devil is in the details, work closely with your vacuum experts in any step of the project, especially design, and ask for vacuum validation tests as soon as possible.

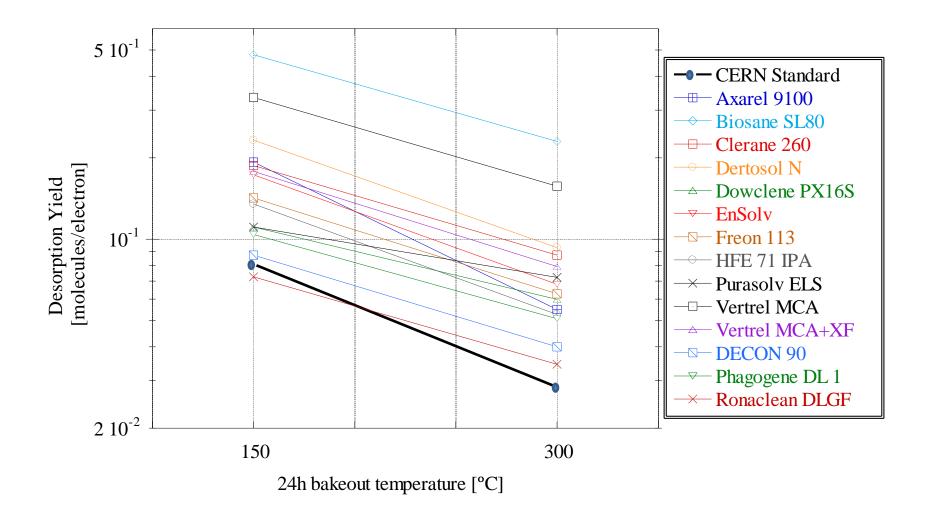
Additional slides

Gas sources: outgassing

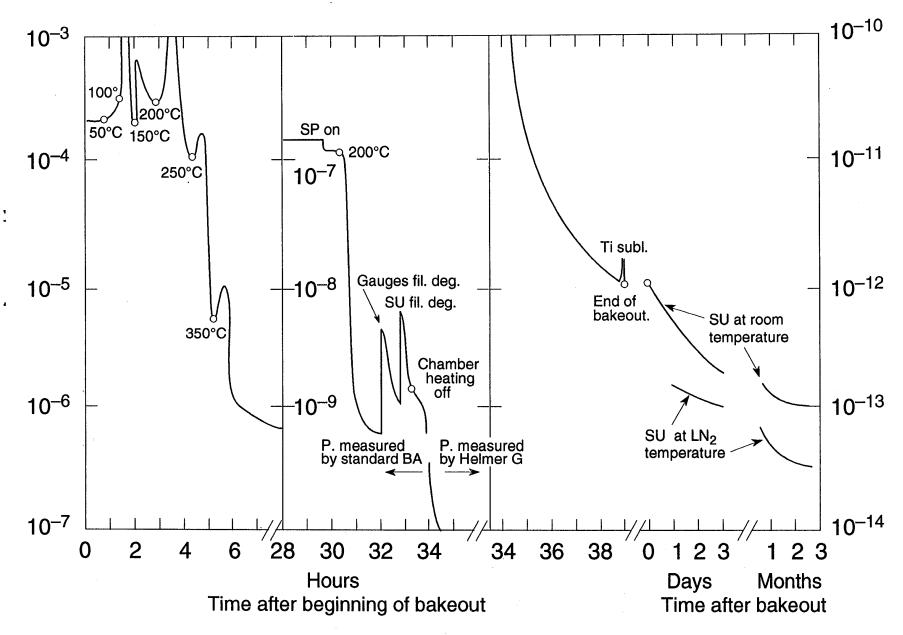


Gas sources: ESD

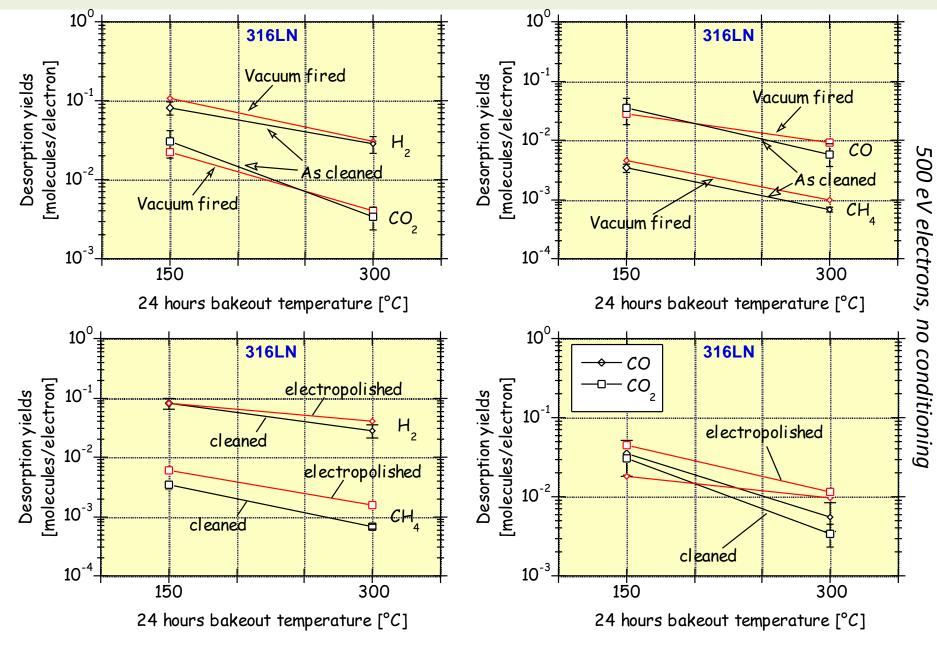
Effect of bakeout temperature and cleaning product

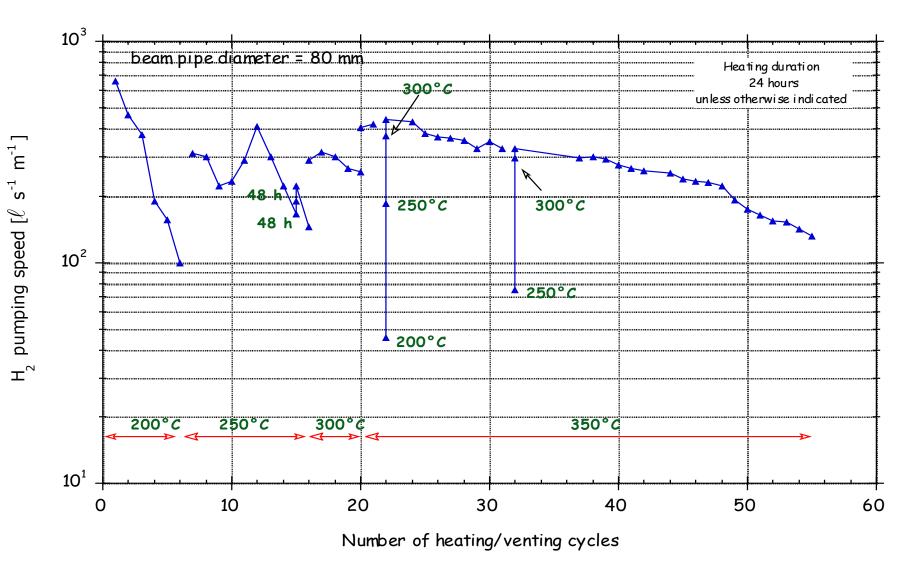


Operational issues



Gas sources: ESD





Cryopumps

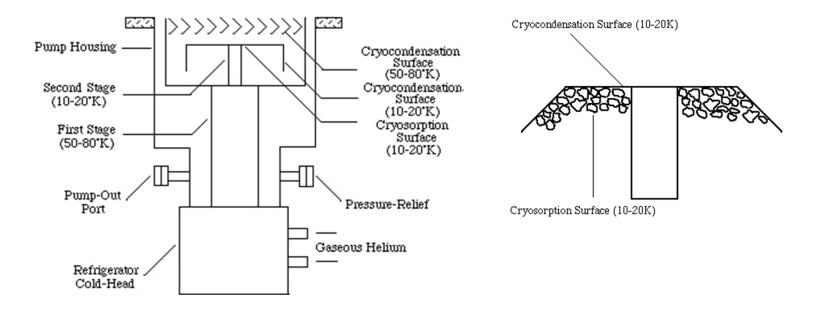


Table 13: Pumping speeds and maximum gas capacities of a commercial

 cryopump (Oerlikon-Leybold 800 BL UHV); the pump inlet diameter is 160 mm.

	Courtesy of Oerlikon-Leybold					
		H ₂ O	N_2	Ar	H_2	He
Very high pumping speeds	S [<i>l</i> s ⁻¹]	2600	800	640	1000	300
	Capacity		225000	225000	3225	375
	[Torr l]		223000	223000		575

Cryosorption and cryocondensation are used for distributed pumping, in particular in vacuum chambers installed in superconducting magnets.