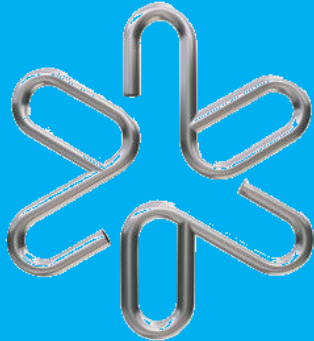


Ciência e Tecnologia do Vácuo

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N. H. Medina

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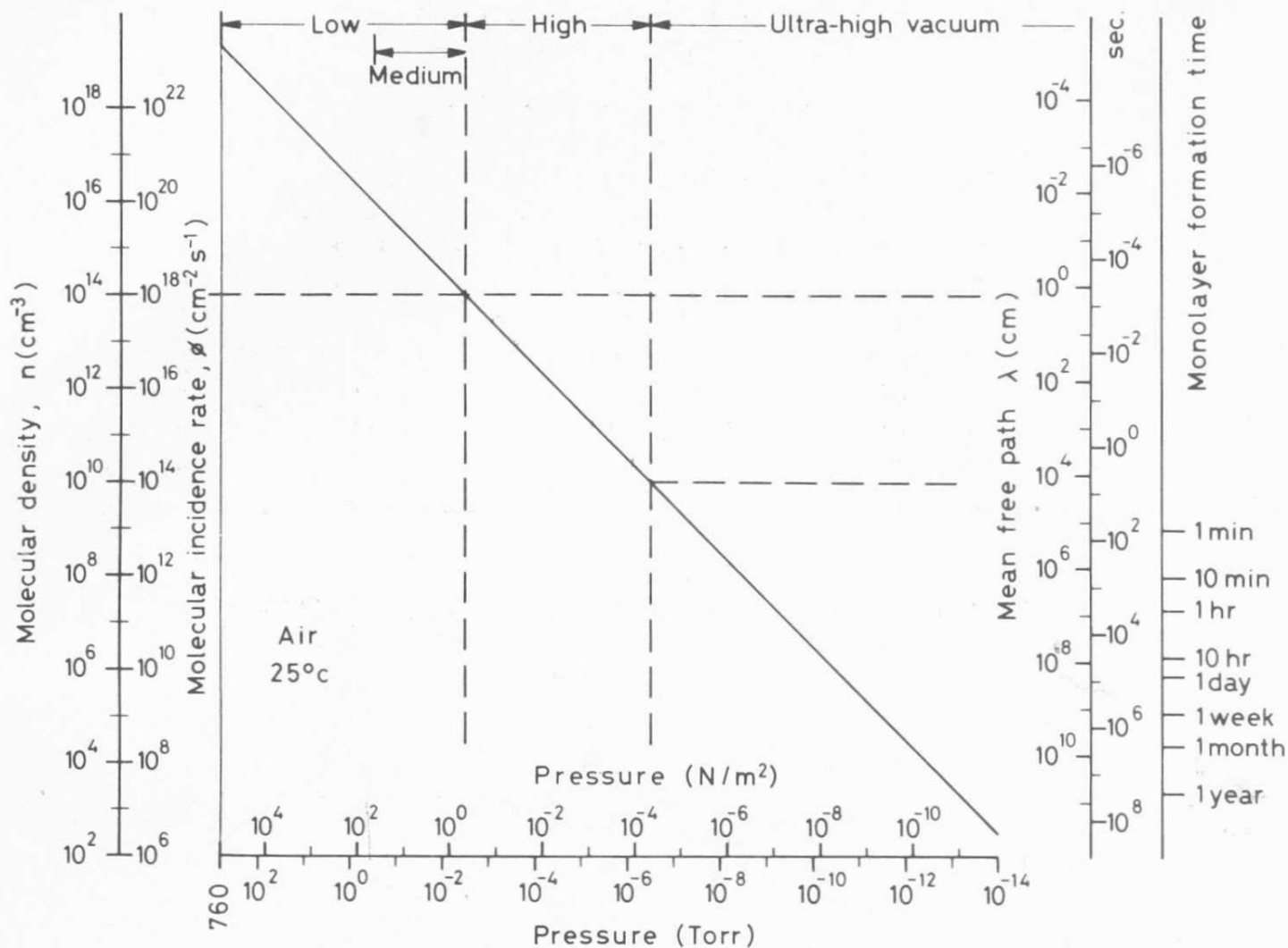
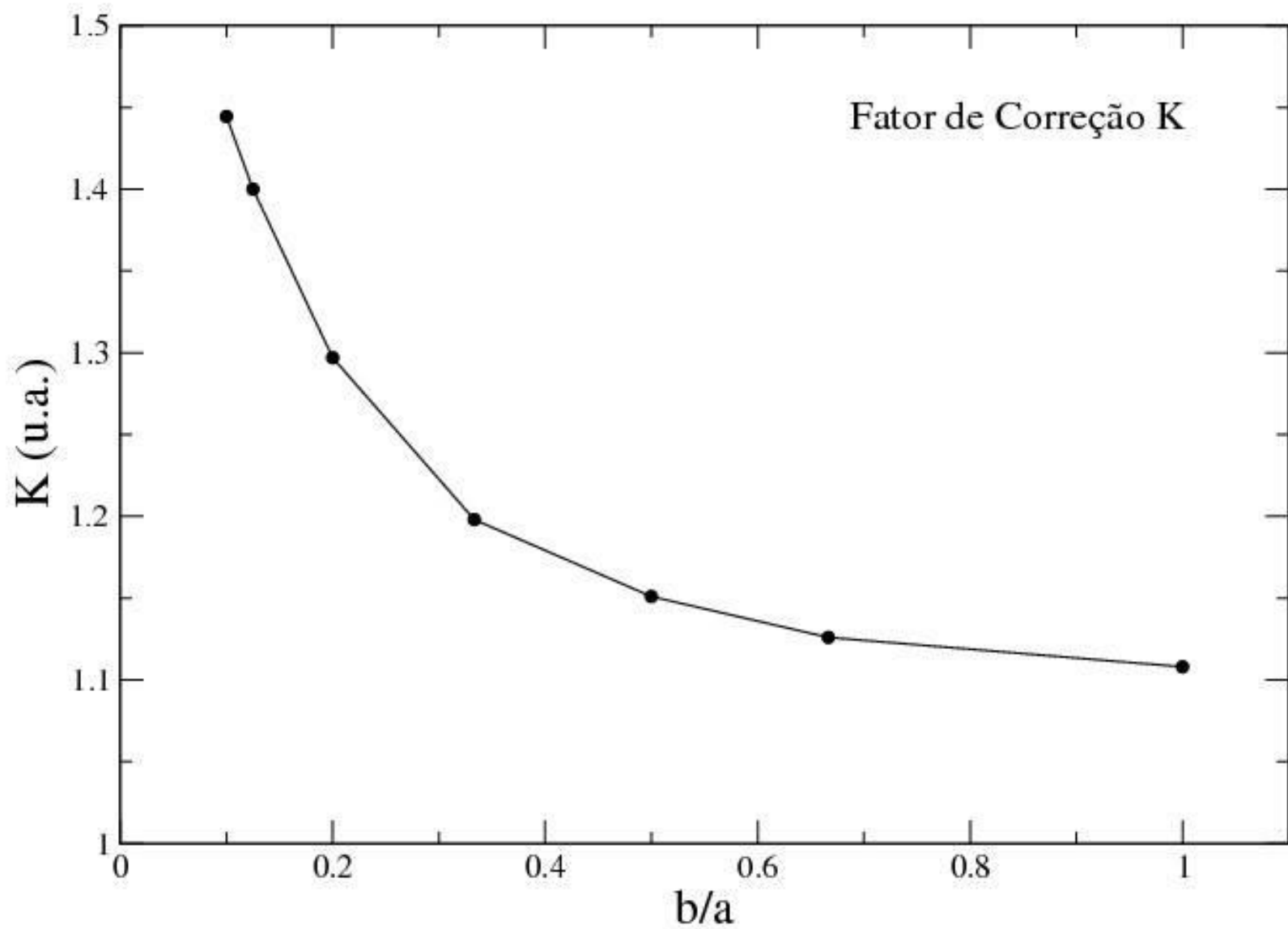


Fig. 1.1 Relationship of several concepts defining the degree of vacuum.

Table 1.3.
Gas compositions.

Component	Atmosphere ⁽¹⁾		Ultra-high vacuum	
	Percent by volume	Partial pressure Torr	(2) Partial pressure Torr	(3)
N ₂	78.08	5.95×10^2	2×10^{-11}	—
O ₂	20.95	1.59×10^2	—	3×10^{-13}
Ar	0.93	7.05	6×10^{-12}	—
CO ₂	0.033	2.5×10^{-1}	6.5×10^{-11}	6×10^{-12}
Ne	1.8×10^{-3}	1.4×10^{-2}	5.2×10^{-11}	—
He	5.24×10^{-4}	4×10^{-3}	3.6×10^{-1}	—
Kr	1.1×10^{-4}	8.4×10^{-4}	—	—
H ₂	5.0×10^{-5}	3.8×10^{-4}	1.79×10^{-9}	2×10^{-11}
Xe	8.7×10^{-6}	6.6×10^{-5}	—	—
H ₂ O	1.57	1.19×10^1	1.25×10^{-10}	9×10^{-13}
CH ₄	2×10^{-4}	1.5×10^{-3}	7.1×10^{-11}	3×10^{-13}
O ₃	7×10^{-6}	5.3×10^{-5}	—	—
N ₂ O	5×10^{-5}	3.8×10^{-4}	—	—
CO	—	—	1.4×10^{-10}	9×10^{-12}

(1) Norton (1962) p. 11, (2) Dennis and Heppel (1968) p. 105, (3) Singleton (1966) p. 355.



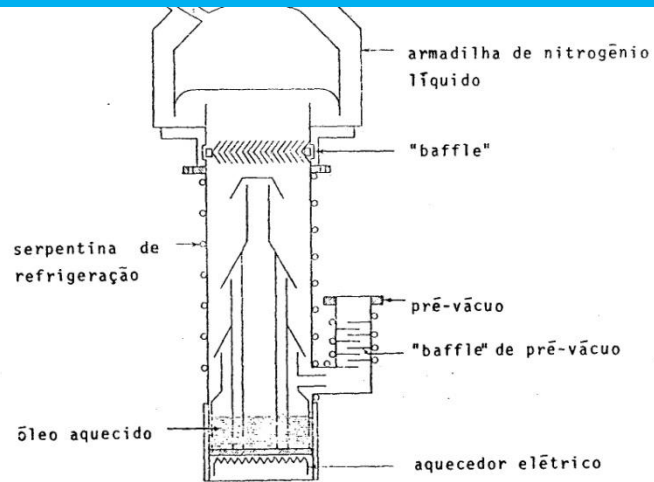


Fig. 29 - Bomba de Difusão com "baffle" chevron e armadilha de nitrogênio líquido ("cold trap")

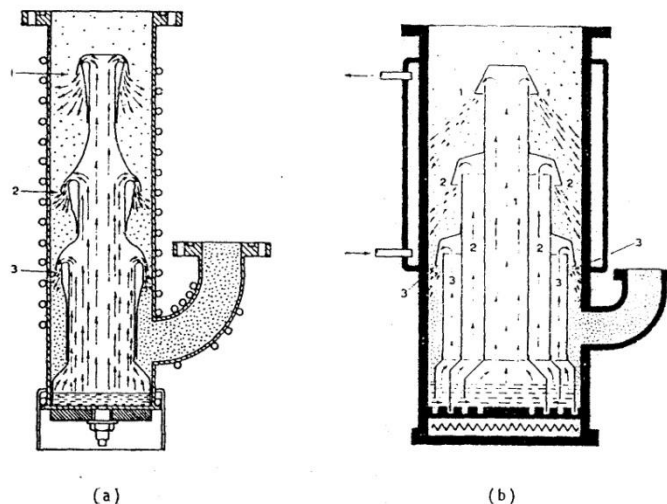


Fig. 30 - Esquemas de Bombas de Difusão de três estágios

- a) desenho mais antigo; o óleo aquecido não sofre nenhum processo de purificação
- b) com tubos concêntricos permitindo a purificação do óleo por destilação fracionada, durante o funcionamento (o vapor de óleo mais aquecido e limpo sai pelo chapéu ("nozzle") 1).

Bomba de Difusão

Jato de vapor empurra as moléculas da câmara criando um gradiente de pressão.

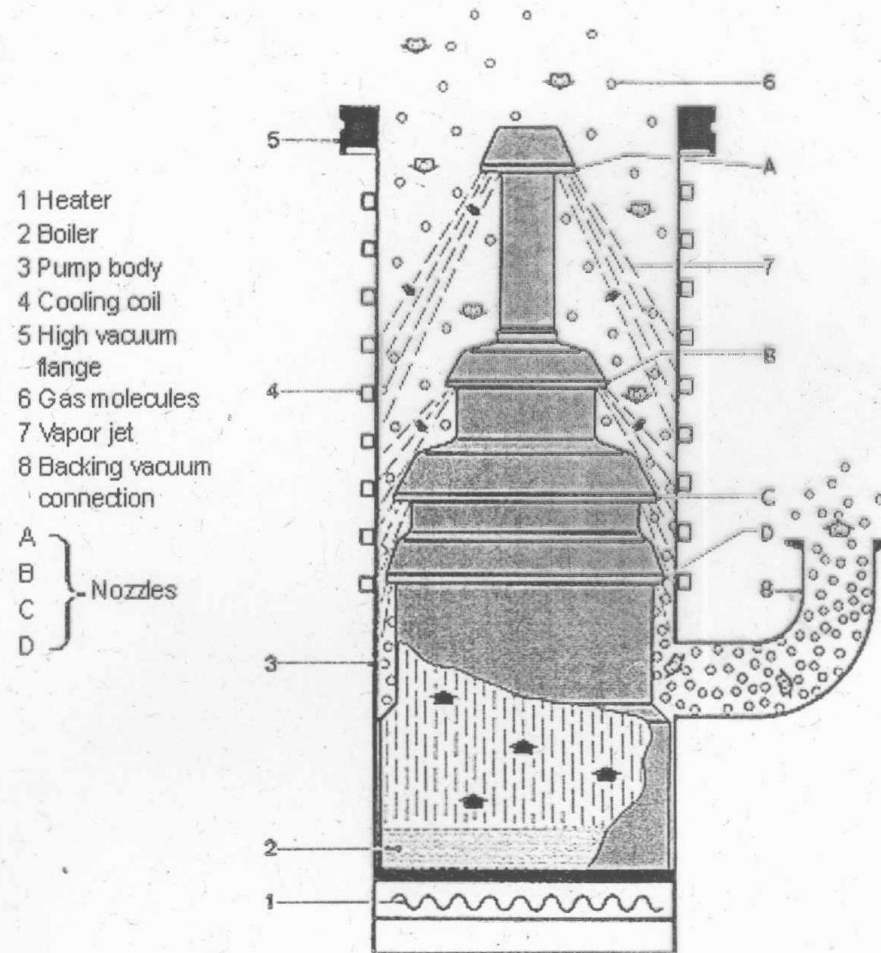


Fig. 2.44
Mode of operation of a diffusion pump

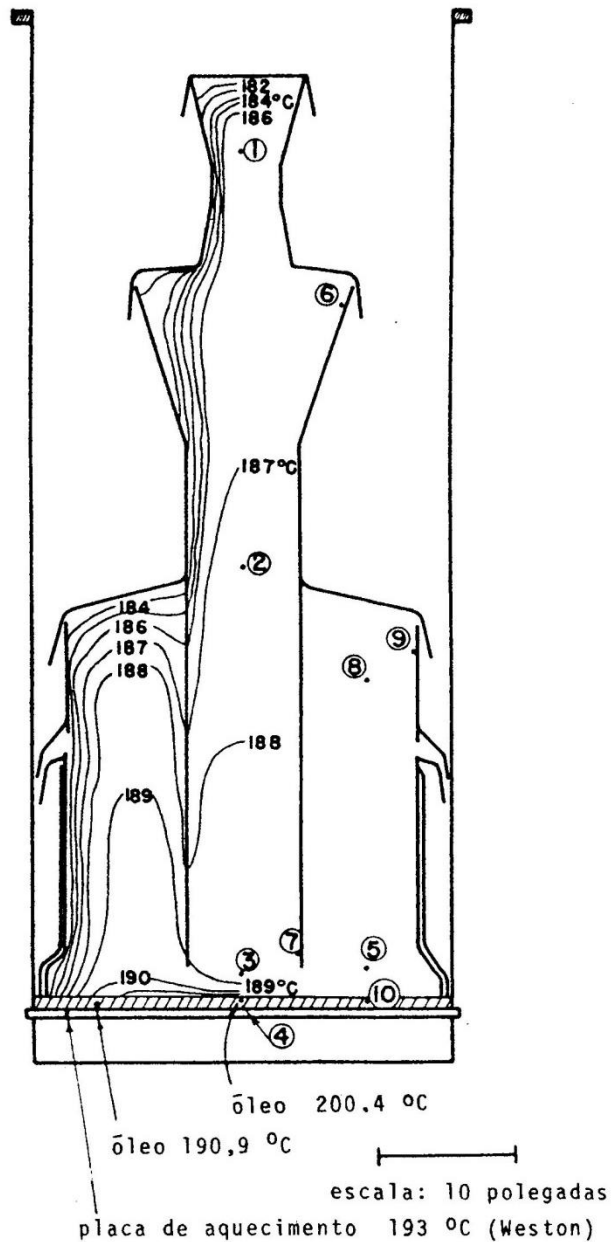
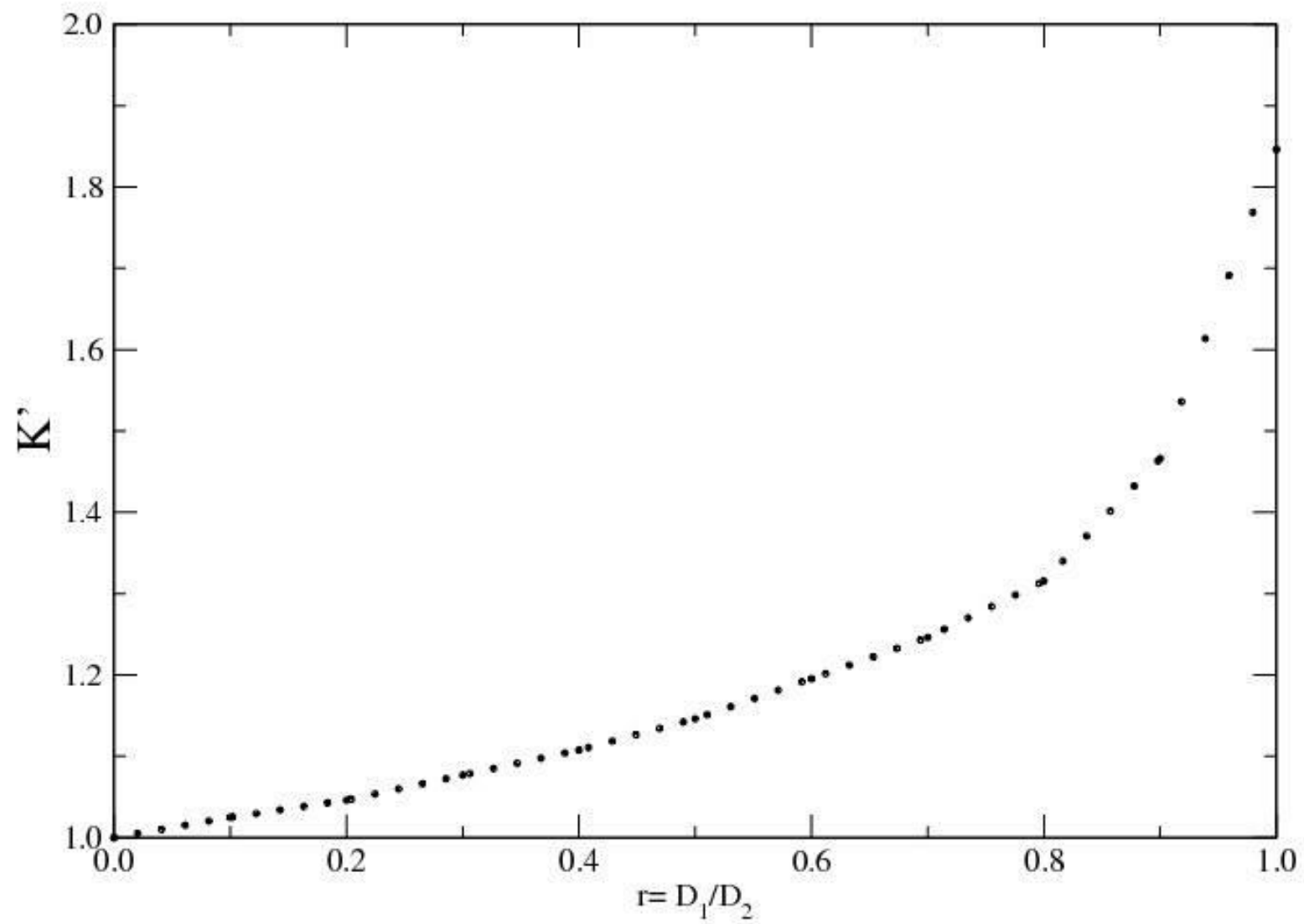
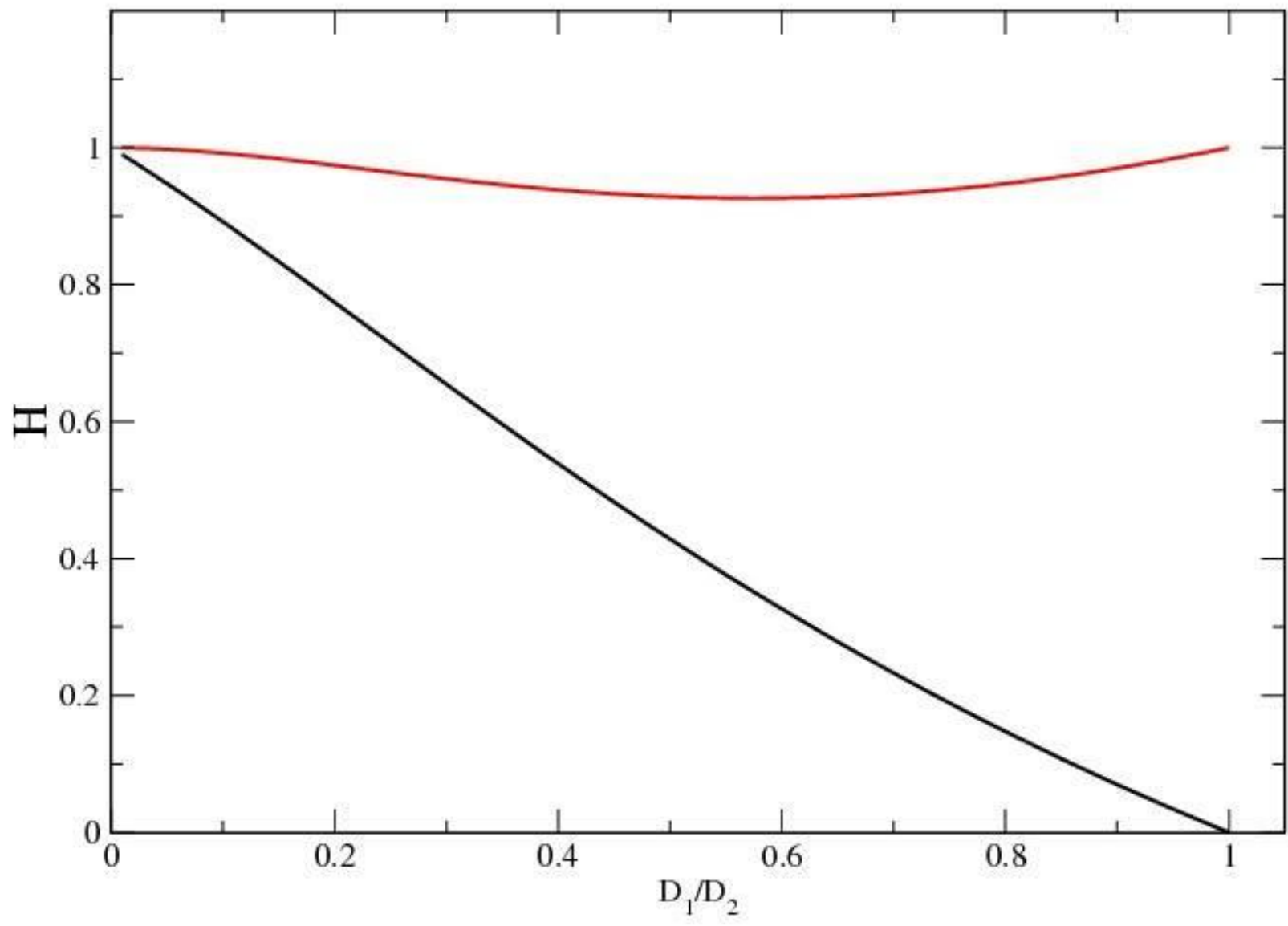


Fig. 31 - Diagrama isotérmico de uma Bomba de Difusão de 32".
 Os pontos numerados representam a localização dos
 termopares utilizados nas medições das temperaturas.





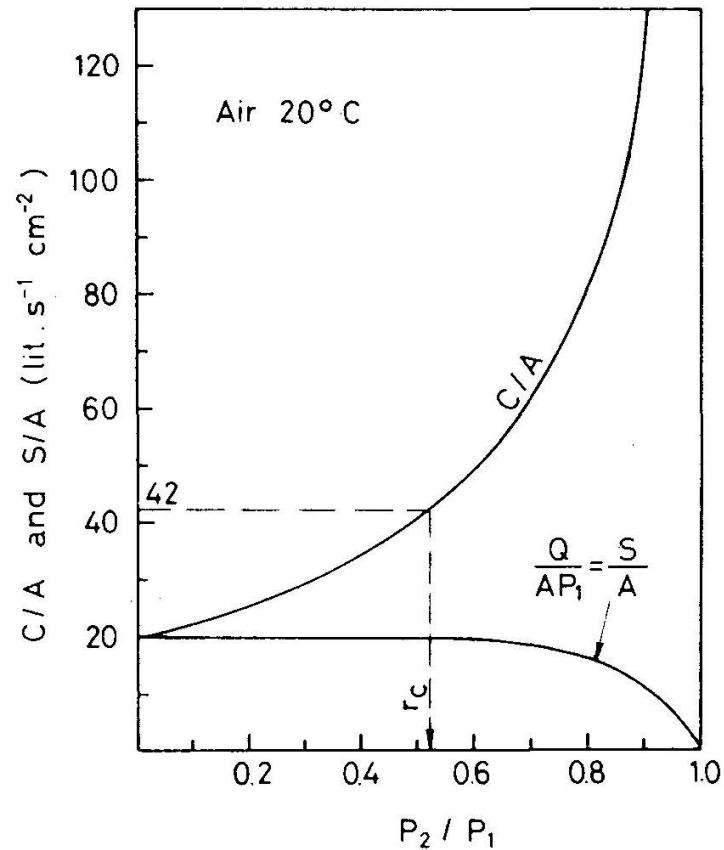


Fig. 3.6 Conductance C and pumping speed S of apertures (viscous flow). A is the cross section area of the aperture.

This equation is plotted for air in fig. 3.36, by using as a parameter the value

$$D^4/L = (128/\pi) \eta E$$

and considering $P_i = 10^6$ dyne/cm² (760 Torr), and $P = 10^2$ dyne/cm² (7.6×10^{-2} Torr), i.e. the pressure range in which usually the flow is viscous. If a volume of $V = 100$ liter is evacuated by a pump of $S_p = 2$ lit/sec through a pipe $D = 2$ cm and $L = 200$ cm, then $D^4/L = 8 \times 10^{-2}$. On the curve 8×10^{-2} , for $S_p = 2$, it results $t/V = 6$ sec/liter. Thus the time required for 100 liter is $t = 600$ sec. If the volume is connected directly to the pump, the line $D^4/L = \infty$ gives $t/V = 4.5$ sec/liter, thus $t = 450$ sec.

It is interesting to mention that if the pump is connected directly to the vessel, $L = 0$, thus $E = \infty$, eq. (3.252) becomes

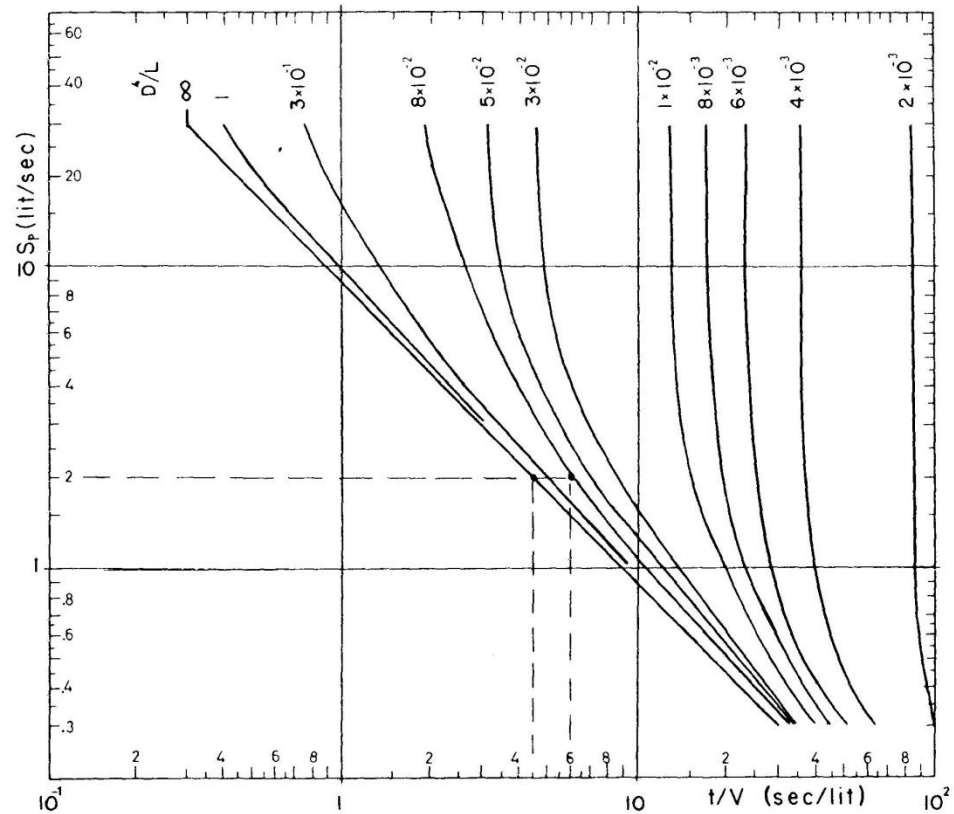


Fig. 3.36 Time required to decrease the pressure from 760 Torr to 7.6×10^{-2} Torr in a volume $V(l)$, connected by a pipe of diameter $D(cm)$ and length $L(cm)$ to a pump of pumping speed $S_p(l/s)$. After Delafosse and Mongodin (1961).

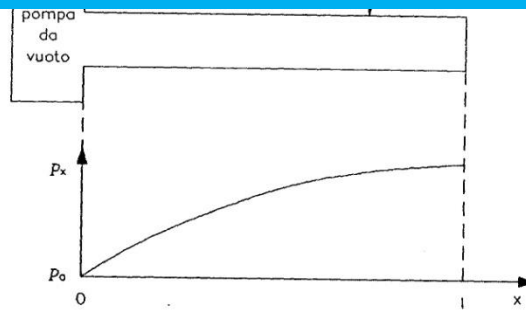


Figura 4.2 - Andamento della pressione in una camera da vuoto tubolare, chiusa ad una estremità e collegata con una pompa all'altra.

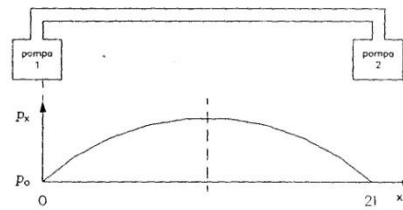


Figura 4.3 - Andamento della pressione in una camera da vuoto tubolare pompata ad entrambe le estremità.

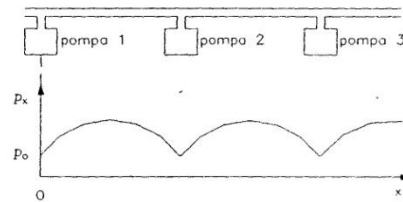


Figura 4.4 - Andamento della pressione in una camera da vuoto tubolare pompata con una serie di pompe disposte ad intervalli regolari di spazio e di uguali caratteristiche.

Bruno Ferrario

Introduzione alla tecnologia
del VUOTO

seconda edizione riveduta ed ampliata da
ANITA CALCATPELLI

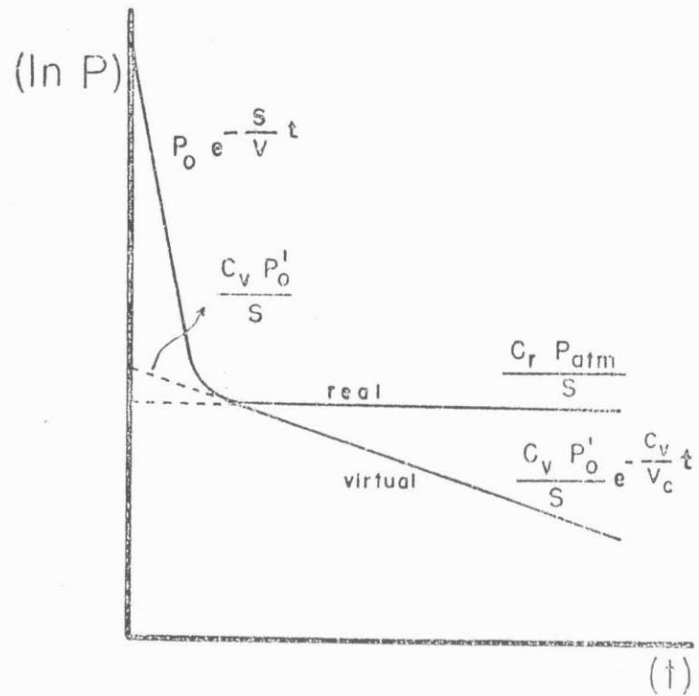


Fig. 2 - Vazamentos: real e virtual.

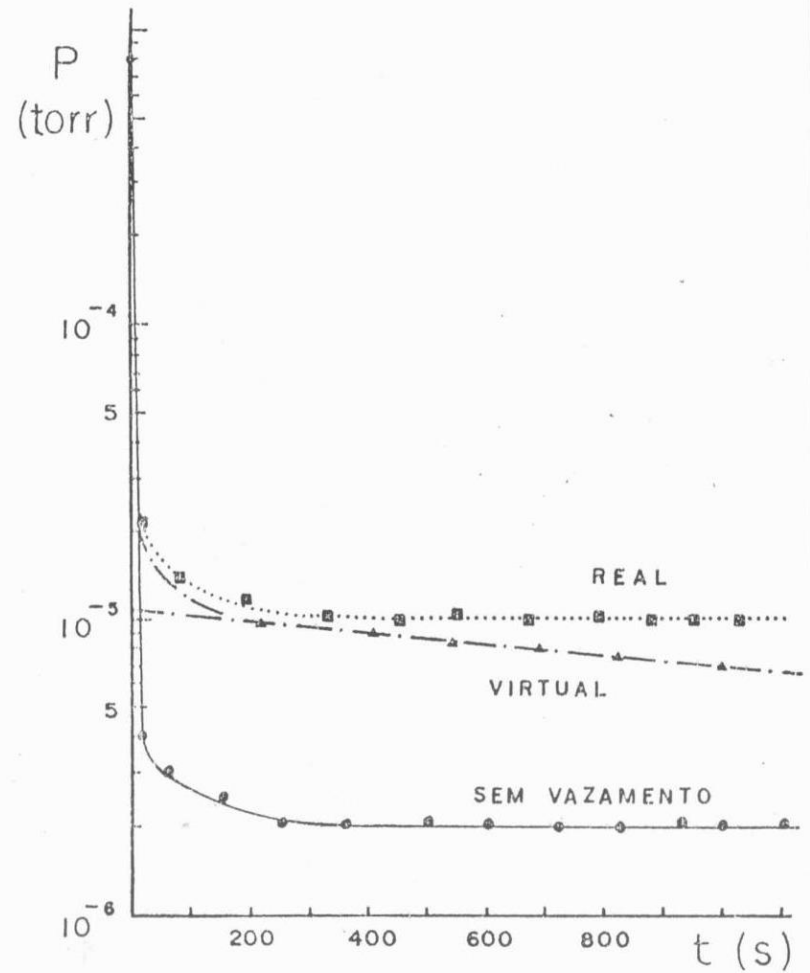
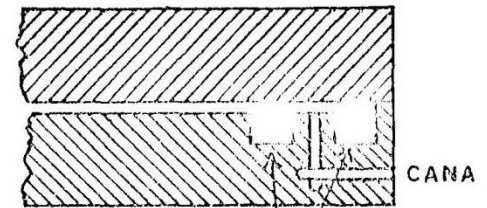
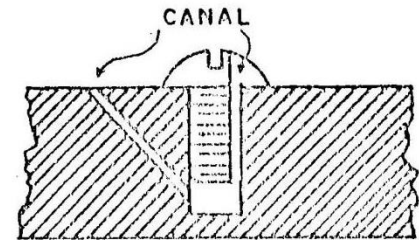
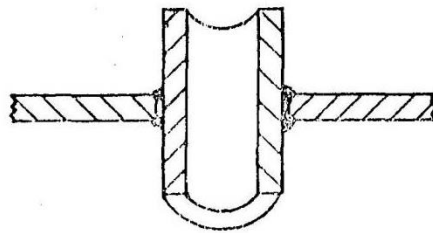
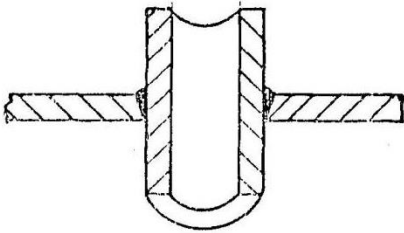
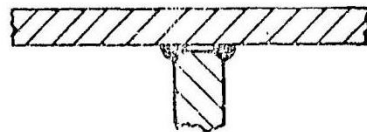
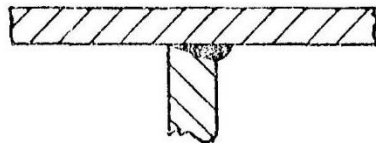
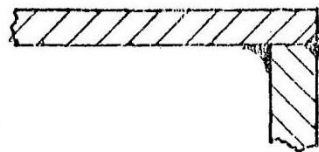
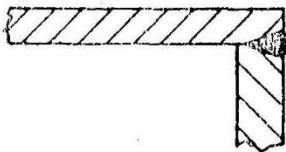
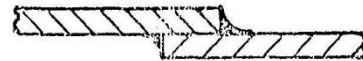


Fig. 5 - Medidas de Simulação de Vazamentos.

CORRETO

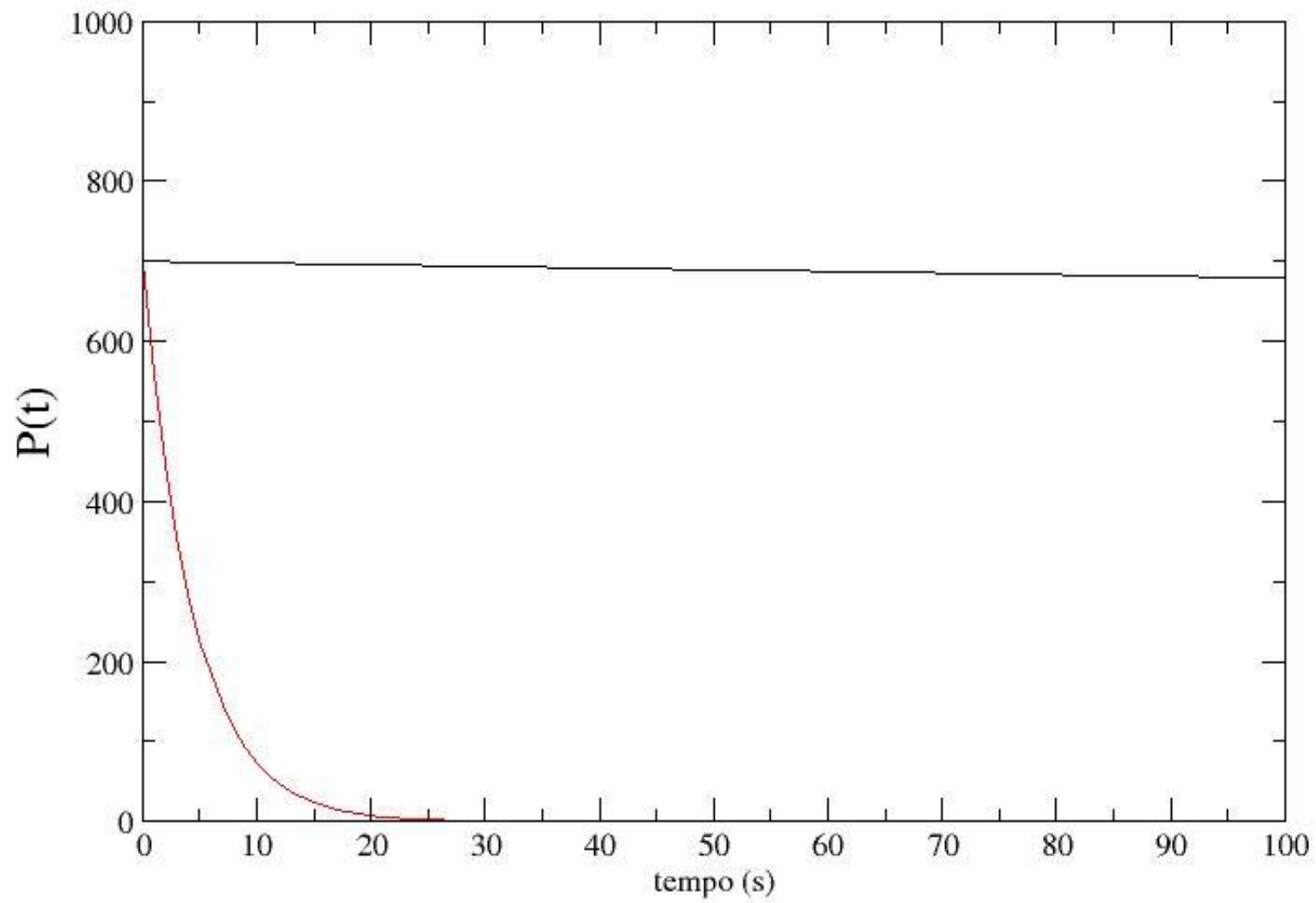
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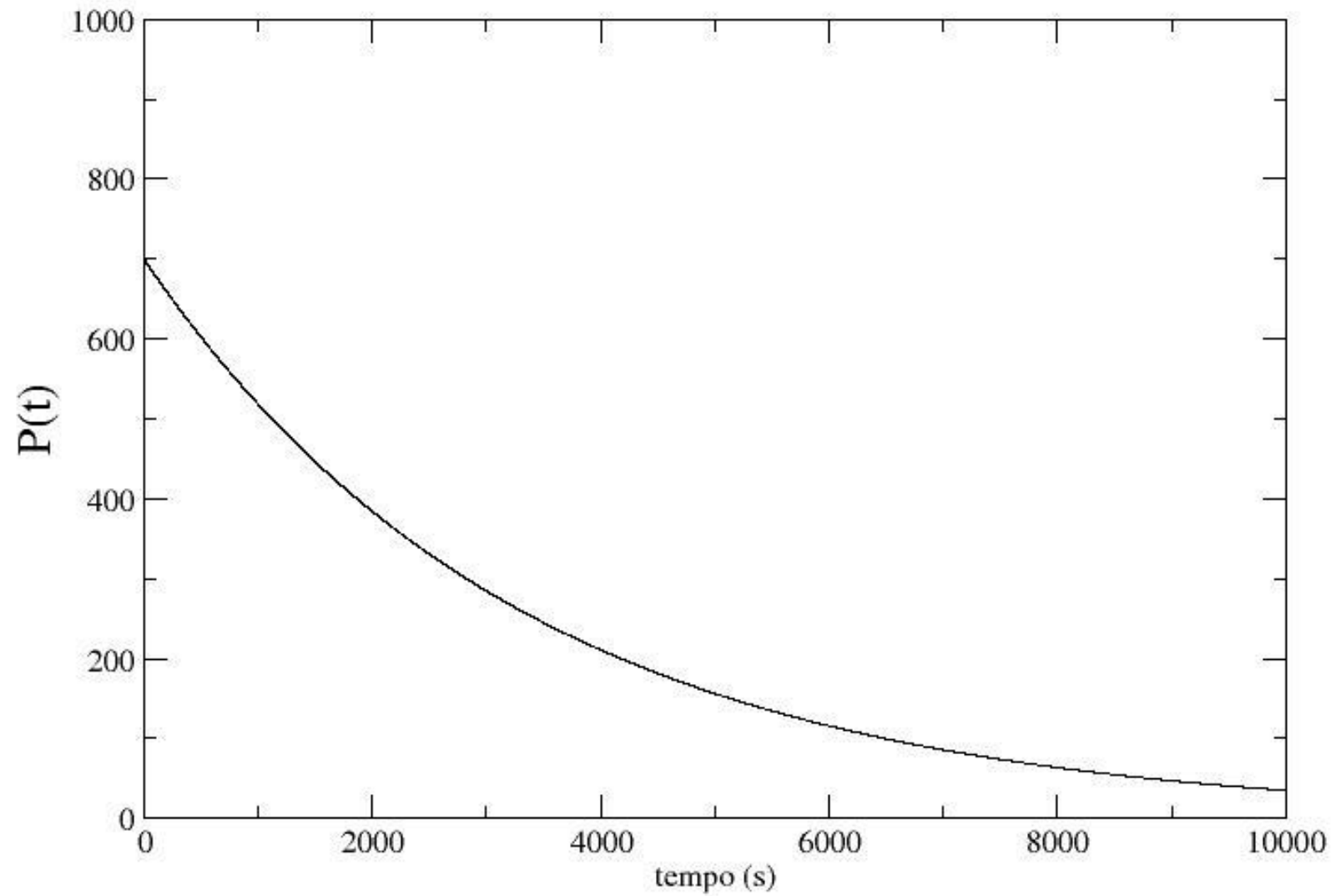
O'RINGS

Fig. 8 - Exemplos de Vazamentos Virtuais e suas correções.

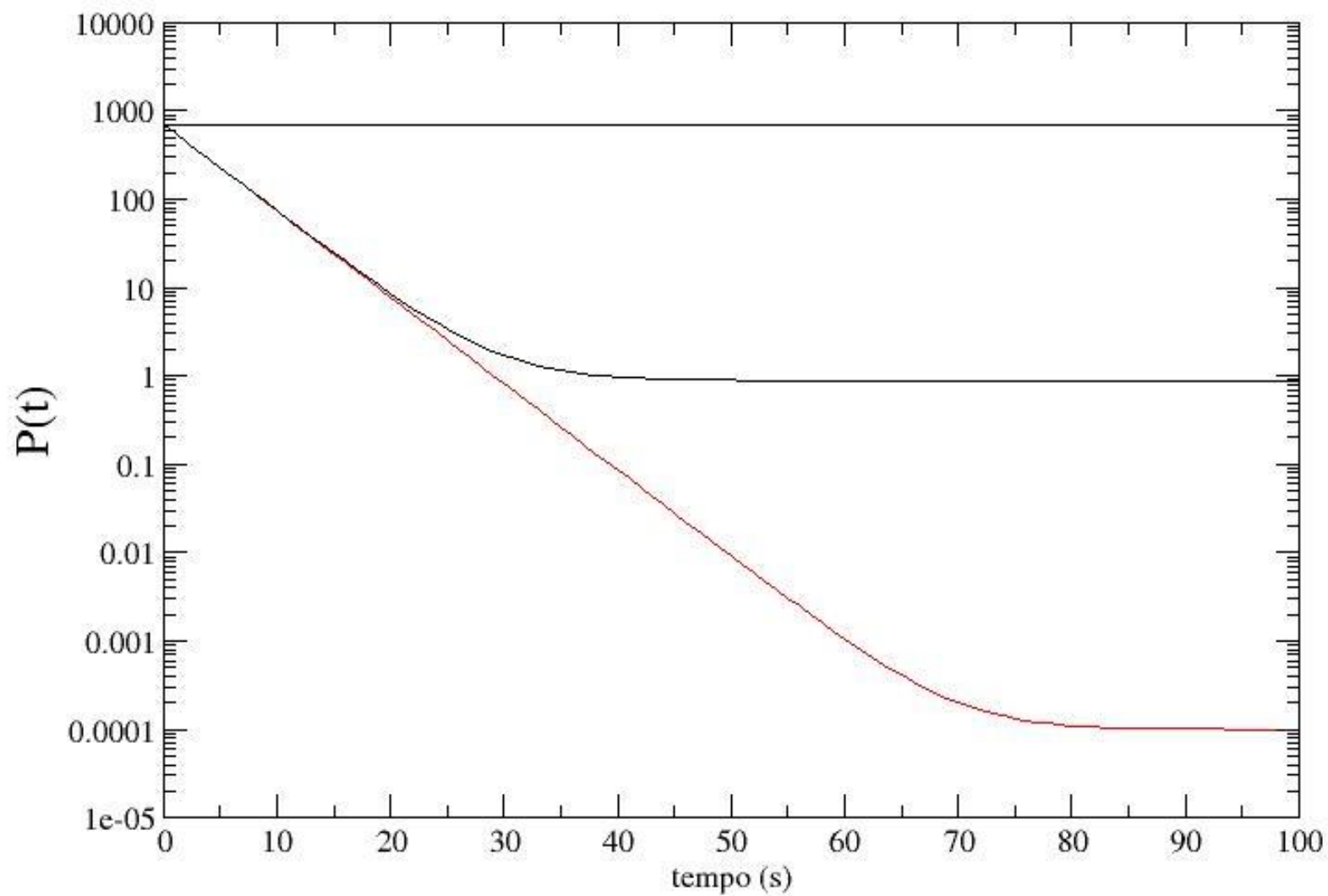
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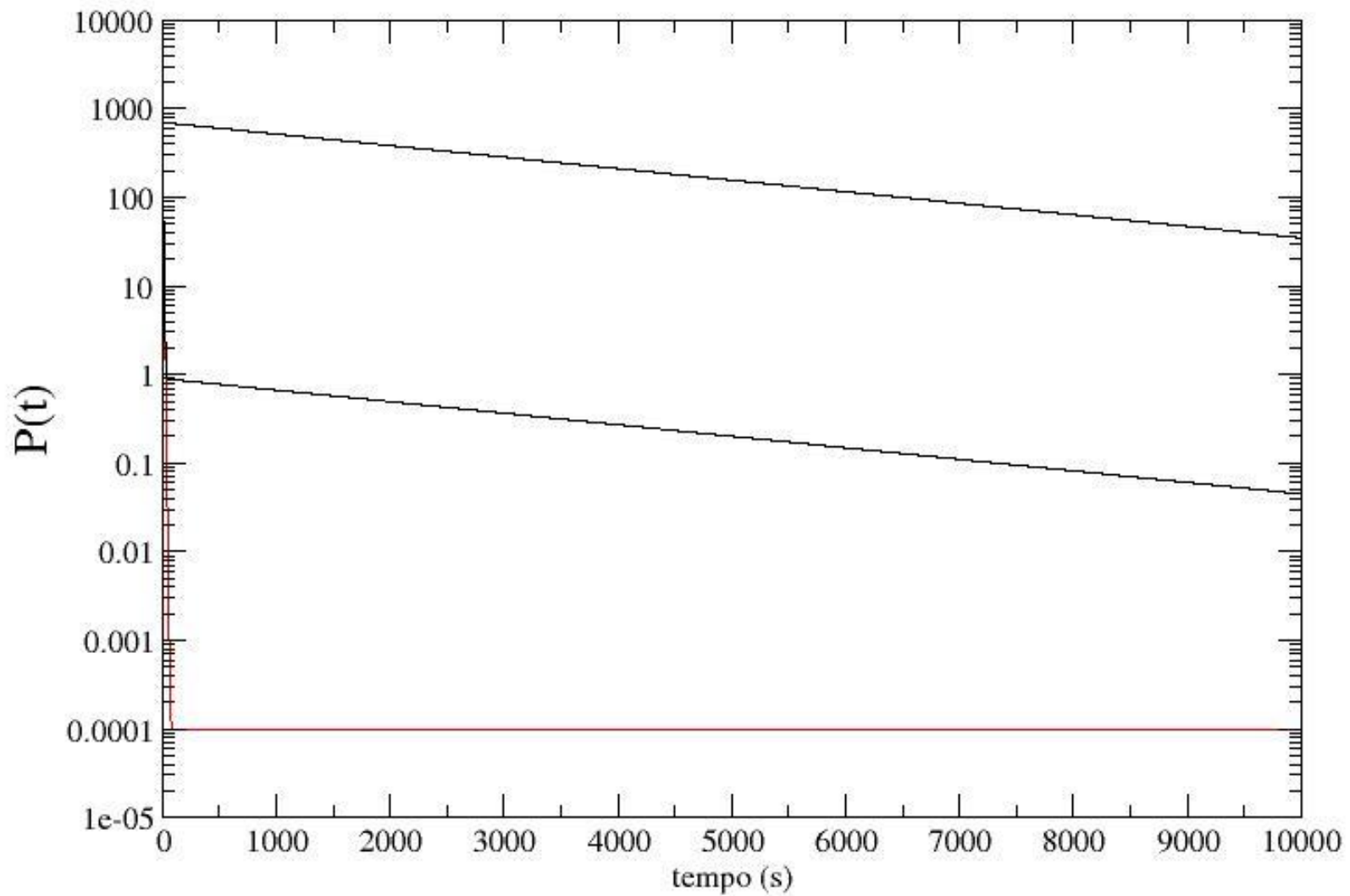
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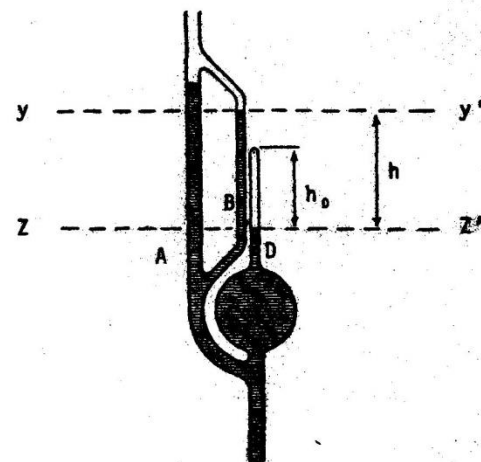
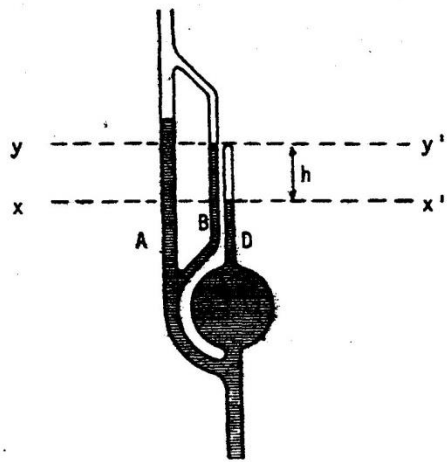
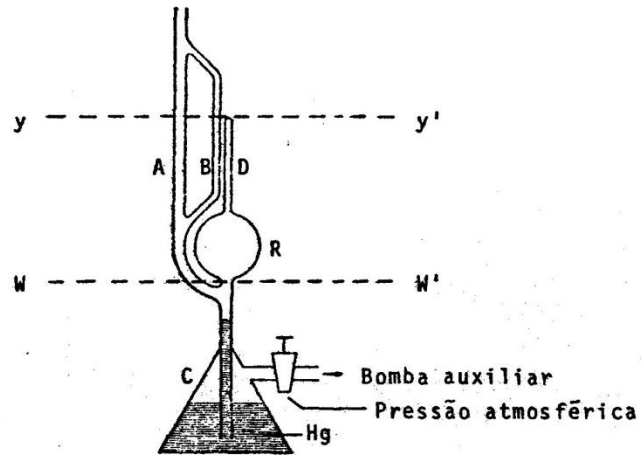


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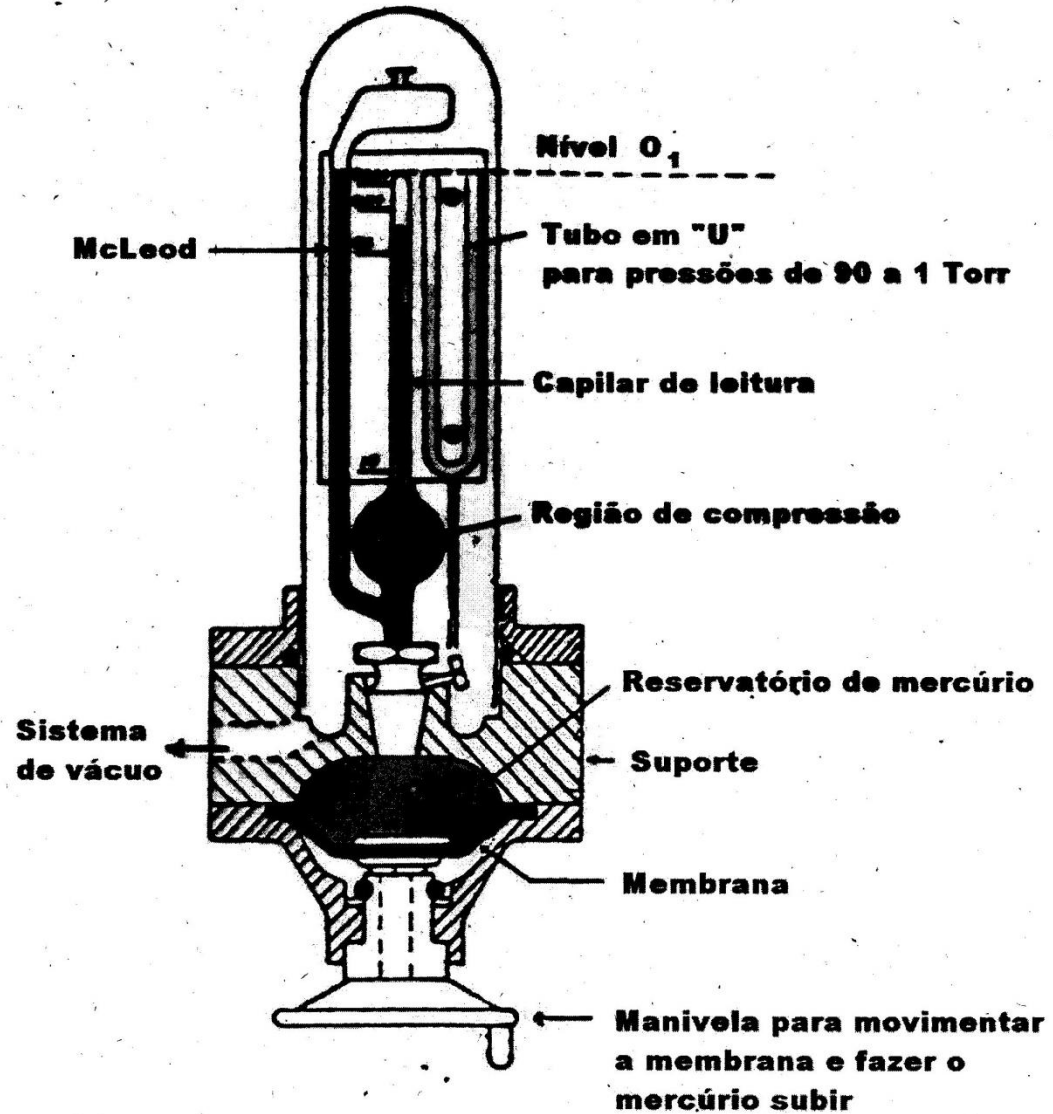


P1P2





Kammerer (McLeod)



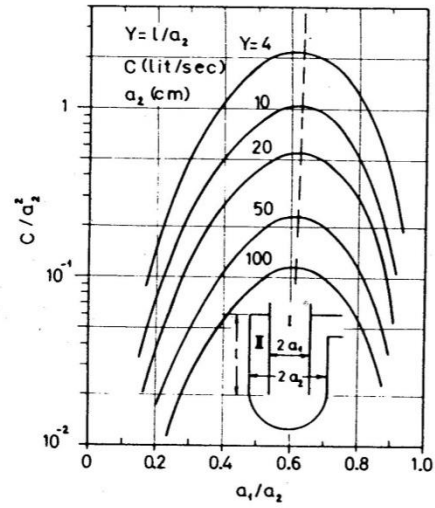


Fig. 3.12 Conductance of traps, for air, 25°C. From Dushman and Lafferty (1962), by permission of J. Wiley & Sons Inc., New York.

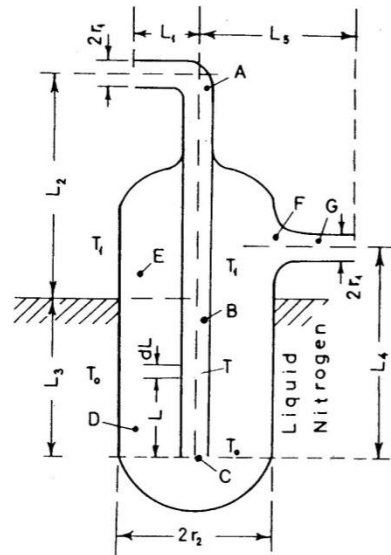
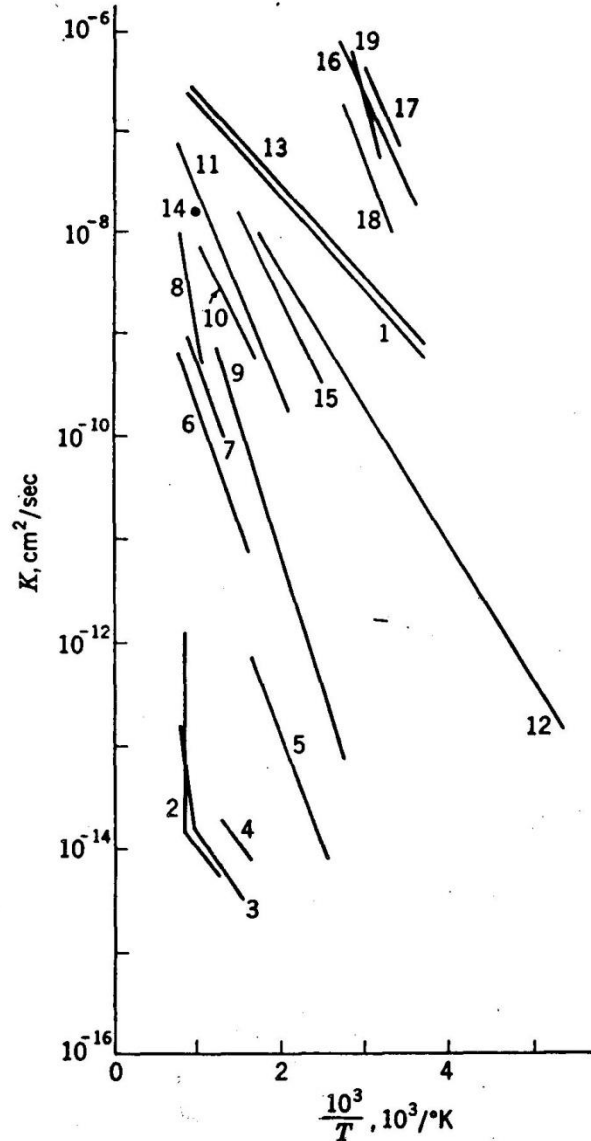


Fig. 3.13 Parts of a liquid nitrogen trap, for calculating its conductance. After Henry (1971).

Fig. 3-2 Permeation constants for various gas-non-metal combinations as a function of temperature. Units are square centimeters per second. [Quantity of gas in cubic centimeters (STP) passing per second through a wall of 1-cm² area and 1-cm thickness, when a pressure difference of 1 atm exists across the wall.] (Numbers 2 to 12 and 14 from P. A. Redhead, J. P. Hobson, and E. V. Kornelsen, *Adv. Electron. Electron Phys.*, vol. 17, p. 323, 1962; Nos. 1, 13, and 15 from V. O. Altemose, *J. Appl. Phys.*, vol. 32, p. 1309, 1961; and 16 to 19 from B. B. Dayton, 1959 *Vacuum Symp. Trans.*, p. 101, 1960.)

1. He-fused silica
2. Air-Pyroceram
3. Air-97% alumina ceramic
4. Air-Pyrex
5. He-lead borate glass G
6. UO₂-97% alumina ceramic
7. Ne-Vycor
8. N₂-SiO₂
9. He-1720 glass
10. He-Pyroceram 9606
11. H₂-SiO₂
12. He-Pyrex 7740
13. He-Vycor 7900
14. H₂-Pyrex
15. He-Pyrex 7052
16. He-Neoprene
17. H₂-Neoprene
18. N₂-Neoprene
19. A-Neoprene



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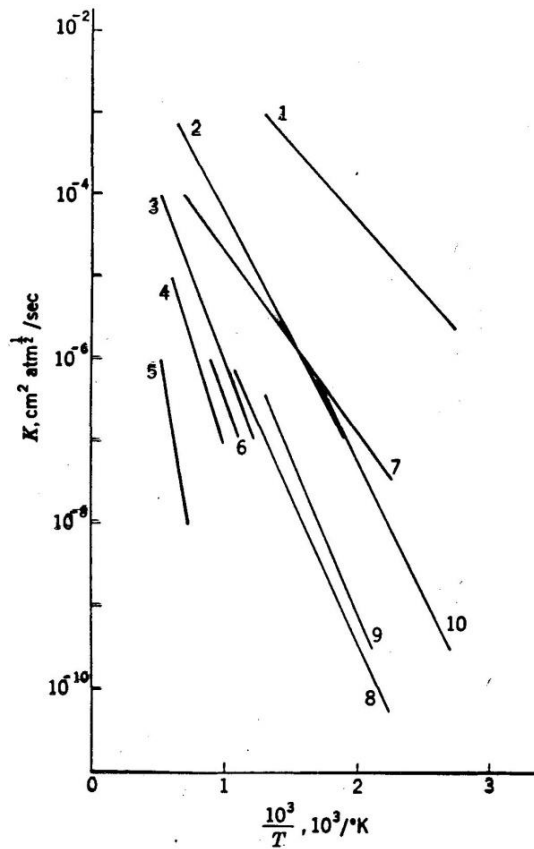


Fig. 3-3 Permeation constants for various diatomic gas-metal combinations as a function of temperature. Units are $\text{cm}^2 \text{ atm}^{1/2} / \text{sec}$. [Quantity of gas in cubic centimeters (STP) passing per second through a wall of 1-cm^2 area and 1-cm thickness, when a pressure difference of 1 atm exists across the wall.] (Numbers 1 to 8 from P. A. Redhead, J. P. Hobson, and E. V. Kornelsen, *Adv. Electron. Electron Phys.*, vol. 17, p. 323, 1962; Nos. 9 and 10 from H. L. Eschbach, F. Gross, and S. Schulien, *Vacuum*, vol. 13, p. 543, 1963.)

- | | |
|---------------------------|--|
| 1. $\text{H}_2\text{-Pd}$ | 6. CO-Fe |
| 2. $\text{H}_2\text{-Ni}$ | 7. $\text{H}_2\text{-Fe}$ |
| 3. $\text{H}_2\text{-Mo}$ | 8. $\text{H}_2\text{-Cu}$ |
| 4. $\text{N}_2\text{-Fe}$ | 9. $\text{H}_2\text{-300 series stainless steel}$ |
| 5. $\text{N}_2\text{-Mo}$ | 10. $\text{H}_2\text{-400 series stainless steel}$ |

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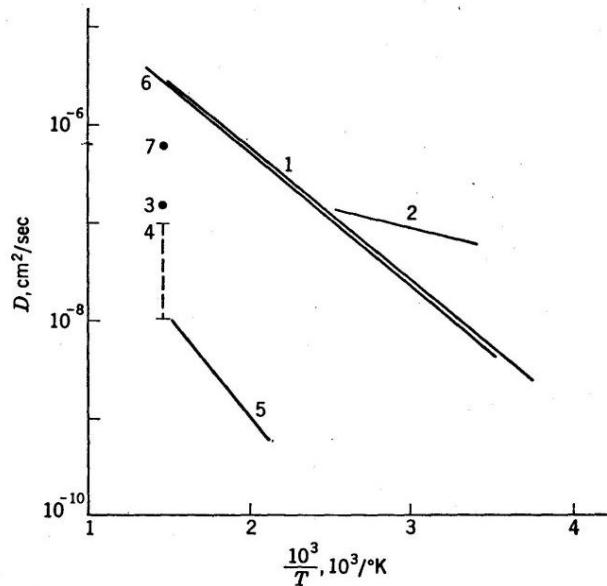


Fig. 3-8 Diffusion constants for various gas-nonmetal combinations. Units are square centimeters per second. (Numbers 1 to 6 from P. A. Redhead, J. P. Hobson, and E. V. Kornelsen, *Adv. Electron. Electron Phys.*, vol. 17, p. 323, 1962, and No. 7 from V. O. Altemose, *J. Appl. Phys.*, vol. 32, p. 1309, 1961.)

- | | |
|--------------------------|-------------------------------------|
| 1. He-Pyrex 7740 | 5. H ₂ -SiO ₂ |
| 2. He-Vycor | 6. He-Duran glass |
| 3. H ₂ -Vycor | 7. He-Pyrex 7052 |
| 4. N ₂ -Vycor | |

Table 3-2 Degassing of a Finite Slab. Error of Approximation [Eq. (3-14)] and fraction of gas removed

Dt/d^2	$\frac{[2Q_T/c_0d]_{\text{Eq. (3-14)}} - [Q_T/c_0d]_{\text{Eq. (3-19)}}}{[Q_T/c_0d]_{\text{Eq. (3-19)}}$	$[Q_T/c_0d]_{\text{Eq. (3-19)}}$
0.06	0.0029	0.55
0.07	0.0057	0.59
0.08	0.014	0.63
0.09	0.016	0.67
0.10	0.023	0.70
0.15	0.072	0.82
0.20	0.14	0.89
0.25	0.21	0.93
0.30	0.29	0.96

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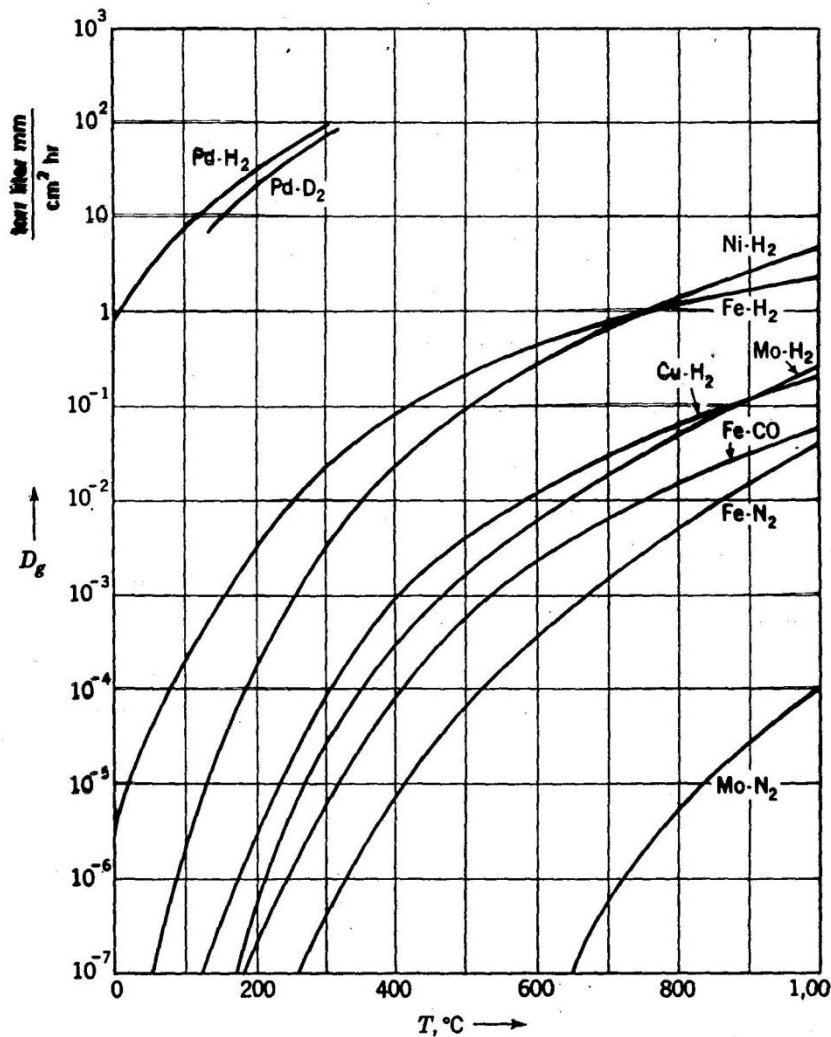


Fig. 3-25 Gas permeation of metal walls versus temperature. Permeation rate D_g is the pressure rise per hour in a vessel of 1-liter volume for a wall area of 1 cm^2 , a wall thickness of 1 mm, and a gas pressure of 760 torr on the outside. T is the temperature of the wall. (From M. von Ardenne, Tabellen Elektronenphysik Ionenphysik und Übermikroskopie, vol. 2, VEB Deutscher Verlag der Wissenschaften, Berlin, 1956.)

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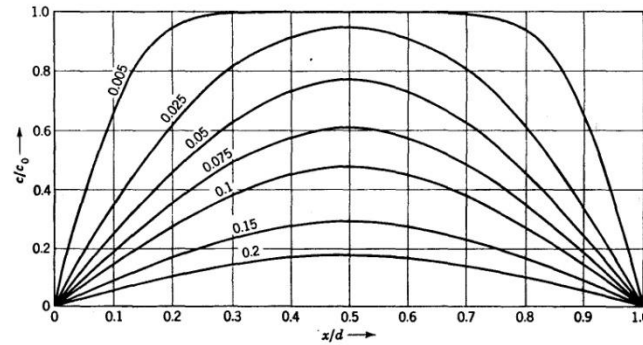


Fig. 3-7 Relative concentration in a finite slab of thickness d for various (dimensionless) times Dt/d^2 as parameter.

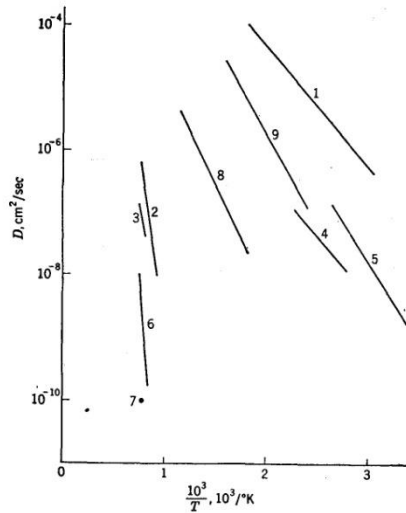


Fig. 3-9 Diffusion constants for various gas-metal combinations. Units are square centimeters per second. (Numbers 1 to 7 from P. A. Redhead, J. P. Hobson, and E. V. Kornelsen, *Adv. Electron. Electron Phys.*, vol. 17, p. 323, 1962; Nos. 8 and 9 from H. L. Eschbach, F. Gross, and S. Schullien, *Vacuum*, vol. 13, p. 543, 1963.)

- | | |
|-----------------------|---|
| 1. H ₂ -Pd | 6. O ₂ -Ni |
| 2. N ₂ -Fe | 7. O ₂ -Fe |
| 3. CO-Ni | 8. H ₂ -300 series stainless steel |
| 4. H ₂ -Ni | 9. H ₂ -400 series stainless steel |
| 5. H ₂ -Fe | |

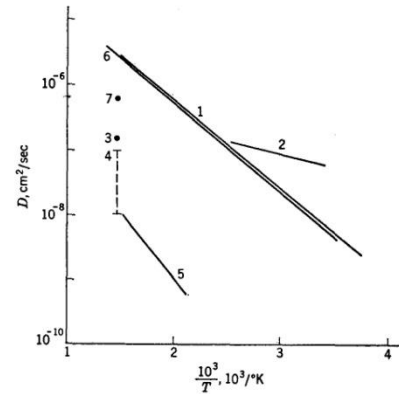


Fig. 3-8 Diffusion constants for various gas-nonmetal combinations. Units are square centimeters per second. (Numbers 1 to 6 from P. A. Redhead, J. P. Hobson, and E. V. Kornelsen, *Adv. Electron. Electron Phys.*, vol. 17, p. 323, 1962, and No. 7 from V. O. Altemose, *J. Appl. Phys.*, vol. 32, p. 1309, 1961.)

- | | |
|--------------------------|-------------------------------------|
| 1. He-Pyrex 7740 | 5. H ₂ -SiO ₂ |
| 2. He-Vycor | 6. He-Duran glass |
| 3. H ₂ -Vycor | 7. He-Pyrex 7052 |
| 4. N ₂ -Vycor | |

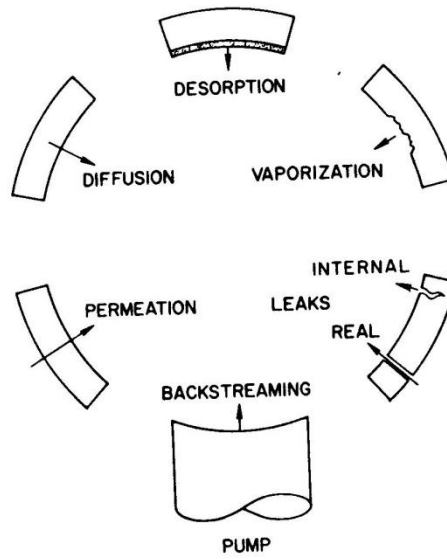


Fig. 4.1 Potential sources of gases and vapors in a vacuum system.

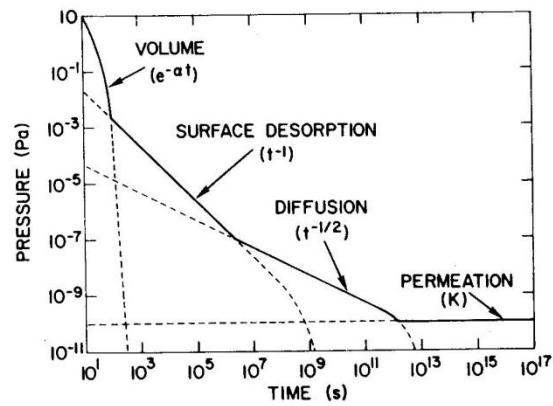


Fig. 4.6 Rate limiting steps during the pumping of a vacuum chamber.

Modelos de Fontes de Gases em um Sistema de Vácuo

Fonte de gás	Característica	Comentário
Volume	$P = P_o e^{-\frac{S}{V}t}$	Pressão cai exponencialmente dependendo de S e V
Vazamento Real	$P_{res} = \frac{Q_{vr}}{S}; Q_{vr} \approx C_{vr} P_{atm}$	Fluxo constante. Utilizar detector de vazamentos. Deve ser eliminado
Vazamento Virtual	$Q_{vv} = \frac{C_v}{S} P_o^n e^{-\frac{C_v}{V_c}t}$	$C_v \ll S_b$ Queda da pressão depende de C_v e V_c . Evitar no projeto
Difusão	$Q(t) = c_o \frac{\sqrt{D}}{\sqrt{\pi t}}$	Q(t) é proporcional a $\frac{1}{\sqrt{t}}$
Permeação	$Q = \frac{K(P_e^n - P_i^n)}{d}$	N=1 para não metais; n=1/2 para moléculas diatômicas em metais. Constante de permeação K(T) é proporcional a $10^3/T$
Evaporação	$W = 0.058 P_v \sqrt{\frac{M}{T}} \frac{g}{cm^2 s}$ $Q = WA \text{ (g/s)}$ $Q = \frac{\Delta N}{\Delta t} kT \frac{\text{Torrl}}{s}$	Crescimento de P_v em função da temperatura é exponencial e por isso mais rápido do que $\frac{1}{\sqrt{T}}$
Desorção Térmica (desgaseificação)	Primeira ordem: $\frac{dc}{dt} = c_o k_1 e^{-\frac{t}{\tau_{res}}}$ $\frac{1}{k_1} = \tau_{res} = \tau_o e^{\frac{E_d}{N k T}}$	Rápido $\tau_{res} = 10^{-12} s$
Temperatura (cozimento)	Segunda ordem: $\frac{dc}{dt} = \frac{-k_2 C_o^2}{(1 + C_o K_2 t)^2}$	Cai lentamente A molécula de H_2 se dissocia na adsorção e recombina na desorção
Superfícies Reais	$q_n = \frac{q}{t^\alpha}$ $0.7 \leq \alpha \leq 2$	Fórmula geral $q_n = q t^{-1}$ Adsorção química Adsorção física

4.6 PRESSURE LIMITS

75

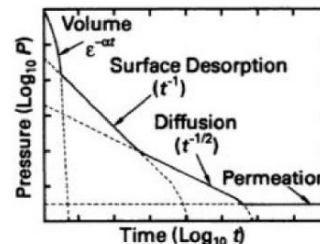


Fig. 4.10 Rate limiting steps during the pumping of a vacuum chamber.

Desorção Térmica

4.2 THERMAL DESORPTION

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Table 4.1 Average Residence Time of Chemisorbed Molecules

System	Desorption Energy (MJ/(kg-mole))	Residence time at		
		77 K (s)	22°C (s)	450°C (s)
H ₂ O/H ₂ O	40.6	10 ¹⁵	10 ⁻⁵	10 ⁻⁹
H ₂ O/metal	96	-	10 ⁵	10 ⁻⁵
H ₂ /Mo	160	-	10 ¹⁷	1

amount of gas is expressed in torr liters at the temperature T :

$$\frac{d(PV)_A}{dt} = 3.64sP \left(\frac{T}{M}\right)^{\frac{1}{2}} \quad \text{torr liters/sec cm}^2 \quad (3-21)$$

Here T is the temperature of the impinging molecules. At pressure P and temperature T , n_A molecules occupy a volume V . The temperature of the surface may be different.

Table 3-3 Chemical Adsorption of Gases on Metals at Room Temperature

Metal	Gas							
	N ₂	H ₂	CO	C ₂ H ₄	C ₂ H ₂	O ₂	CO ₂	CH ₄
Ag	N	N	N, Y*					
Al	N	N	Y	Y	Y	Y		
Au	N	N	N, Y*	Y	Y	N		
Ba	Y	Y	Y	Y	Y	Y	Y	Y
Ca	Y	Y	Y	Y	Y	Y		
Cd	N	N	N	N	N	Y		
Co	N	Y	Y			Y		N
Cu	N	N	Y	Y	Y	Y		
Cr	N	Y	Y	Y	Y	Y		Y
Fe	Y	Y	Y	Y	Y	Y		N
Hg	N	N	N					
In	N	N	N	N	N	Y		
Mg			Y					
Mn						Y		
Mo	Y	Y	Y	Y	Y	Y		Y
Nb	Y	Y	Y			Y		
Ni	N	Y	Y	Y	Y	Y		N
Pb	N	N	N	N	N	Y		
Pd	N	Y	Y	Y	Y	Y		Y
Pt	N	Y	Y	Y	Y	Y		
Rh	N	Y	Y	Y	Y	Y		Y
Sn	N	N	N	N	N	Y		
Sr			Y					
Ta	Y	Y	Y	Y	Y	Y		Y
Ti	Y	Y	Y	Y	Y	Y	Y	Y
W	Y	Y	Y	Y	Y	Y		Y
Zn	N	N	N	N	N	Y		
Zr	Y	Y	Y	Y	Y	Y		

N = No, Y = Yes.

* Two different investigators.

SOURCE: J. P. Hobson, *Brit. J. Appl. Phys.*, vol. 14, p. 544, 1963.

Table 3-5 Heats of Adsorption in Kilocalories per mole for Dilute Layers

Chemisorption:			
Rb on W	60	O ₂ on Ni	115
Cs on W	64	H ₂ on Fe	32
B on W	140	N ₂ on Fe	40
Ni on Mo	48	H ₂ on Ir	26
Ag on Mo	35	H ₂ on Rh	26
H ₂ on W	46	H ₂ on Co	~24
O ₂ on W	194	H ₂ on Pt	27
CO on W	~100	O ₂ on Pt	67
N ₂ on W	85	H ₂ on Pd	27
CO ₂ on W	122	H ₂ on Ni	30
NH ₃ on W	70	NH ₃ on Ni	36
H ₂ on Mo	~40	CO on Ni	35
H ₂ on Ta	46	H ₂ on Cu	8
Physical adsorption:			
Xe on W	8-9	Xe on Mo	~8
Kr on W	~4.5	Xe on Ta	~5.3
A on W	~1.9		

SOURCE: G. Ehrlich, *Ann. N.Y. Acad. Sci.*, vol. 101, art. 3, p. 722, 1963, 1961 *Vacuum Sump. Trans.*, D. 126, 1962.

Table 3-4 Activation Energy for Surface Migration E_m and Energy of Desorption E_D in Dilute Adlayers

	E_m E_D		E_m/E_D
	kcal/mole		
Cs on W	14	64	0.22
W on W (110)	30	134	0.22
Ba on W (100)	15	87	0.17
O on W	30	147	0.22
H on W	16	74	0.22
N on W	35	155	0.23
CO on W	65	~100	0.7
Xe on W	3.8	9	0.4
Kr on W	>1.1	4.5	>0.25
A on W	0.6	1.9	0.3
H on Ni	7	67	0.11

SOURCE: G. Ehrlich, *Ann. N.Y. Acad. Sci.*, vol. 101, art. 3, p. 722, 1963.

Appendix A.3 (Continued)

Conventional unit	→ multiply by →	to get SI unit
Gas flow		
micron-L/s	0.13332	Pa-L/s
Pa-L/s	3.6	Pa-m ³ /h
atm-cc/s	101.323	Pa-L/s
Torr-L/s	133.32	Pa-L/s
Torr-L/s	0.133	J/s
watt	1000	Pa-L/s
kg-mole/s (at 0°C)	2.48 × 10 ⁹	Pa-L/s
molecules/s (at 0°C)	4 × 10 ⁻¹⁸	Pa-L/s
Outgassing rate		
Pa-L/(m ² -s)	0.001	W/m ²
Pa-m ³ /(m ² -s)	1.0	W/m ²
μL/(cm ² -s)	1.33	W/m ²
Torr-L/(cm ² -s)	1333.2	W/m ²
Dynamic viscosity		
poise	10	Pa-s
Newton-s/m ²	1	Pa-s
Kinematic viscosity		
centistoke	1	mm ² /s
Diffusion constant		
cm ² /s	0.0001	m ² /s
Heat conductivity		
watt-cm ⁻¹ -K ⁻¹	100	J-s ⁻¹ -m ⁻¹ -K ⁻¹
Specific Heat		
cal-(g-mole) ⁻¹ -K ⁻¹	4184.	J-(kg-mole) ⁻¹ -K ⁻¹
J-kg ⁻¹ -K ⁻¹	<i>M</i>	J-(kg-mole) ⁻¹ -K ⁻¹
BTU-lb ⁻¹ -°F ⁻¹	4186 <i>M</i>	J-(kg-mole) ⁻¹ -K ⁻¹
Heat capacity		
cal-(g-mole) ⁻¹	4184	J-(kg-mole) ⁻¹
J/kg	<i>M</i>	J-(kg-mole) ⁻¹
BTU/lb	2325.9 <i>M</i>	J-(kg-mole) ⁻¹
Energy, work, or quantity of heat		
kW-h	3.6	MJ
kcal	4184	J
BTU	1055	J
ft-lb	1.356	J
to get Conventional unit ←	divide by ←	SI unit

APPENDIX C

Material Properties

Appendix C.1 Outgassing Rates of Vacuum Baked Metals

Material	Treatment	q (10^{-11} W/m ²)
Aluminum ^a	15 h at 250°C	53.0
Aluminum ^b	20 h at 100°C	5.3
6061 Aluminum ^c	glow disch. + 200°C bake	1.3
Copper ^b	20 h at 100°C	146.0
304 Stainless Steel ^a	30 h at 250°C	400.0
Stainless Steel ^d	2 h at 850/900°C vac. furnace	27.0
316L Stainless Steel ^e	2 h at 800°C vac. furnace	46.0
U15C Stainless Steel ^f	3 h vac. furn. 1000°C + 25-h <i>in situ</i> vac. bake at 360°C	2.1

Source. Adapted with permission from *Vacuum*, **25**, p. 347, R. J. Elsey. Copyright 1975, Pergamon Press.

^a J. R. Young, *J. Vac. Sci. Technol.*, **6**, 398 (1969);

^b G. Moraw, *Vacuum*, **24**, 125 (1974);

^c H. J. Halama and J. C. Herrera, *J. Vac. Sci. Technol.*, **13**, 463 (1976);

^d R. L. Samuel, *Vacuum*, **20**, 295 (1970);

^e R. Nuvolone, *J. Vac. Sci. Technol.*, **14**, 1210 (1977);

^f R. Calder and G. Lewin, *Brit. J. Appl. Phys.*, **18**, 1459 (1967).

Appendix C.2 Outgassing Rates of Unbaked Metals¹

Material	q_1 (10^{-7} W/m ²)	α_1	q_{10} (10^{-7} W/m ²)	α_{10}
Aluminum (fresh) ^a	84.0	1.0	8.0	1.0
Aluminum (degassed 24-h) ^a	55.2	3.2	4.08	0.9
Aluminum (3-h in air) ^a	88.6	1.9	6.33	0.9
Aluminum (fresh) ^a	82.6	1.0	4.33	0.9
Aluminum (anodized 2 μ m pores) ^a	3679.0	0.9	429.0	0.9
Aluminum (bright rolled) ^b	-	-	100.0	1.0
Duraluminum ^b	2266.0	0.75	467.0	0.75
Brass (wave guide) ^b	5332.0	2.0	133.0	1.2
Copper (fresh) ^a	533.0	1.0	55.3	1.0
Copper (mech. polished) ^a	46.7	1.0	4.75	1.0
OHFC copper (fresh) ^a	251.0	1.3	16.8	1.3
OHFC copper (mech. polished) ^a	25.0	1.1	2.17	1.1
Gold (wire fresh) ^a	2105.0	2.1	6.8	1.0
Mild steel ^b	7200.0	1.0	667.0	1.0
Mild steel (slightly rusty) ^b	8000.0	3.1	173.0	1.0
Mild steel (chromium plated polished) ^b	133.0	1.0	12.0	-
Mild steel (aluminum spray coated) ^b	800.0	0.75	133.0	0.75
Steel (chromium plated fresh) ^a	94.0	1.0	7.7	1.0
Steel (chromium plated polished) ^a	121.0	1.0	10.7	1.0
Steel (nickel plated fresh) ^a	56.5	0.9	6.6	0.9
Steel (nickel plated) ^a	368.0	1.1	3.11	1.1
Steel (chemically nickel plated fresh) ^a	111.0	1.0	9.4	1.0
Steel (chemically nickel plated polished) ^a	69.6	1.0	6.13	1.0
Steel (descaled) ^a	4093.0	0.6	3933.0	0.7
Molybdenum ^a	69.0	1.0	4.89	1.0
Stainless steel EN58B (AISI 321) ^b	-	-	19.0	1.6
Stainless steel 19/9/1-electropolished ^c	-	-	2.7	-
-vapor degreased ^c	-	-	1.3	-
-Diversey cleaned ^c	-	-	4.0	-
Stainless steel ^b	2333.0	1.1	280.0	0.75
Stainless steel ^b	1200.0	0.7	267.0	0.75
Stainless steel ICN 472 (fresh) ^a	180.0	0.9	19.6	0.9
Stainless steel ICN 472 (sanded) ^a	110.0	1.2	13.9	0.8
Stainless steel NS22S (mech. polished) ^a	22.8	0.5	6.1	0.7
Stainless steel NS22S (electropolished) ^a	57.0	1.0	5.7	1.0
Stainless steel ^a	192.0	1.3	18.0	1.9
Zinc ^a	2946.0	1.4	429.0	0.8
Titanium ^a	150.0	0.6	24.5	1.1
Titanium ^a	53.0	1.0	4.91	1.0

Source. Reprinted with permission from *Vacuum*, 25, p 347, R. J. Elsey. Copyright 1975, Pergamon Press.

¹ $q_n = qt^{-\alpha_n}$, where n is in hours.

^a A. Schram, *Le Vide*, No. 103, 55 (1963),

^b B. B. Dayton, *Trans. 6th Nat. Vac. Symp. (1959)*, Pergamon Press, New York, 1960, p. 101,

^c R. S. Barton and R. P. Govier, *Proc. 4th Int. Vac. Congr. (1968)*, Institute of Physics and the Physical Society, London, 1969, p. 775, and *Vacuum*, 20, 1 (1970).

Appendix C.3 Outgassing Rates of Ceramics and Glasses¹

Material	q_1 (10^{-7} W/m ²)	α_1	q_{10} (10^{-7} W/m ²)	α_{10}
Steatite ^a	1200.0	1.0	127.0	-
Pyrophyllite ^b	2667.0	1.0	267.0	-
Pyrex (fresh) ^c	98.0	1.1	7.3	-
Pyrex (1 month in air) ^c	15.5	0.9	2.1	-

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¹ $q_n = qt^{-\alpha_n}$, where n is in hours.

^a R. Geller, *Le Vide*, No. 13, 71 (1958);

^b R. Jaeckel and F. Schittko, quoted by Elsey;

^c B. B. Dayton, *Trans. 6th Nat. Symp. Vac. Technol. (1959)*, Pergamon Press, New York, 1960, p. 101.

Appendix C.4 Outgassing Rates of Elastomers¹

Material	q_1 (10^{-5} W/m ²)	α_1	q_4 (10^{-5} W/m ²)	α_4
Butyl DR41 ^a	200.0	0.68	53.0	0.64
Neoprene ^a	4000.0	0.4	2400.0	0.4
Perbunan ^a	467.0	0.3	293.0	0.5
Silicone ^b	930.0	-	267.0	-
Viton A (fresh) ^c	152.0	0.8	-	-
Viton A (bake 12 h at 200°C) ^d	-	-	0.027 ^e	-
Polyimide (bake 12 h at 300°C) ^d	-	-	0.005 ^e	-

Source. Adapted with permission from *Vacuum*, **25**, p. 347, R. J. Elsey. Copyright 1975, Pergamon Press.

¹ $q_n = qt^{-\alpha_n}$, where n is in hours.

^a J. Blears, E. J. Greer and J. Nightengale, *Adv. Vac. Sci. Technol.*, **2**, E. Thomas, Ed., Pergamon Press, 1960, p. 473;

^b D. J. Santeler, et al., *Vacuum Technology and Space Simulation*, NASA SP-105, National Aeronautics and Space Administration, Washington, DC, 1966, p. 219;

^c A. Schram, *Le Vide*, No. 103, 55 (1963);

^d P. Hait, *Vacuum*, **17**, 547 (1967);

^e Pumping time is 12 h.

Appendix C.7 Outgassing Rates of Polymers¹

Material	q_1 (10^{-5} W/m ²)	α_1	q_{10} (10^{-5} W/m ²)	α_{10}
Araldite (molded) ^a	155.0	0.8	47.0	0.8
Araldite D ^b	253.0	0.3	167.0	0.5
Araldite F ^b	200.0	0.5	97.0	0.5
Kel-F ^c	5.0	0.57	2.3	0.53
Methyl Methacrylate ^d	560.0	0.9	187.0	0.57
Mylar (24-h at 95 RH) ^e	307.0	0.75	53.0	-
Nylon ^f	1600.0	0.5	800.0	0.5
Plexiglas ^g	961.0	0.44	36.0	0.44
Plexiglas ^b	413.0	0.4	240.0	0.4
Polyester-glass Laminate ^c	333.0	0.84	107.0	0.81
Polystyrene ^c	2667.0	1.6	267.0	1.6
PTFE ^h	40.0	0.45	26.0	0.56
PVC (24-h at 95 RH) ^e	113.0	1.0	2.7	-
Teflon ^g	8.7	0.5	3.3	0.2

Source. Reprinted with permission from *Vacuum*, 25, p. 347, R. J. Elsey, Copyright 1975, Pergamon Press.

¹ $q_n = qt^{-\alpha_n}$, where n is in hours.

^a A. Schram, *Le Vide*, No. 103, 55 (1963);

^b R. Geller, *Le Vide*, No.13, 71 (1958);

^c B. B. Dayton, CVC Technical Report;

^d J. Blears, E. J. Greer and J. Nightengale, *Adv. Vac. Sci. Technol.*, 2, E. Thomas, Ed., Pergamon Press, 1960, p. 473;

^e D. J. Santeler, *Trans. 5th Symp. Vac. Tech. (1958)*, Pergamon Press, New York, 1959, p. 1;

^f B. D. Power and D. J. Crawley, *Adv. Vac. Sci. Technol.*, 1, E. Thomas, Ed., Pergamon Press, New York, 1960, p. 207;

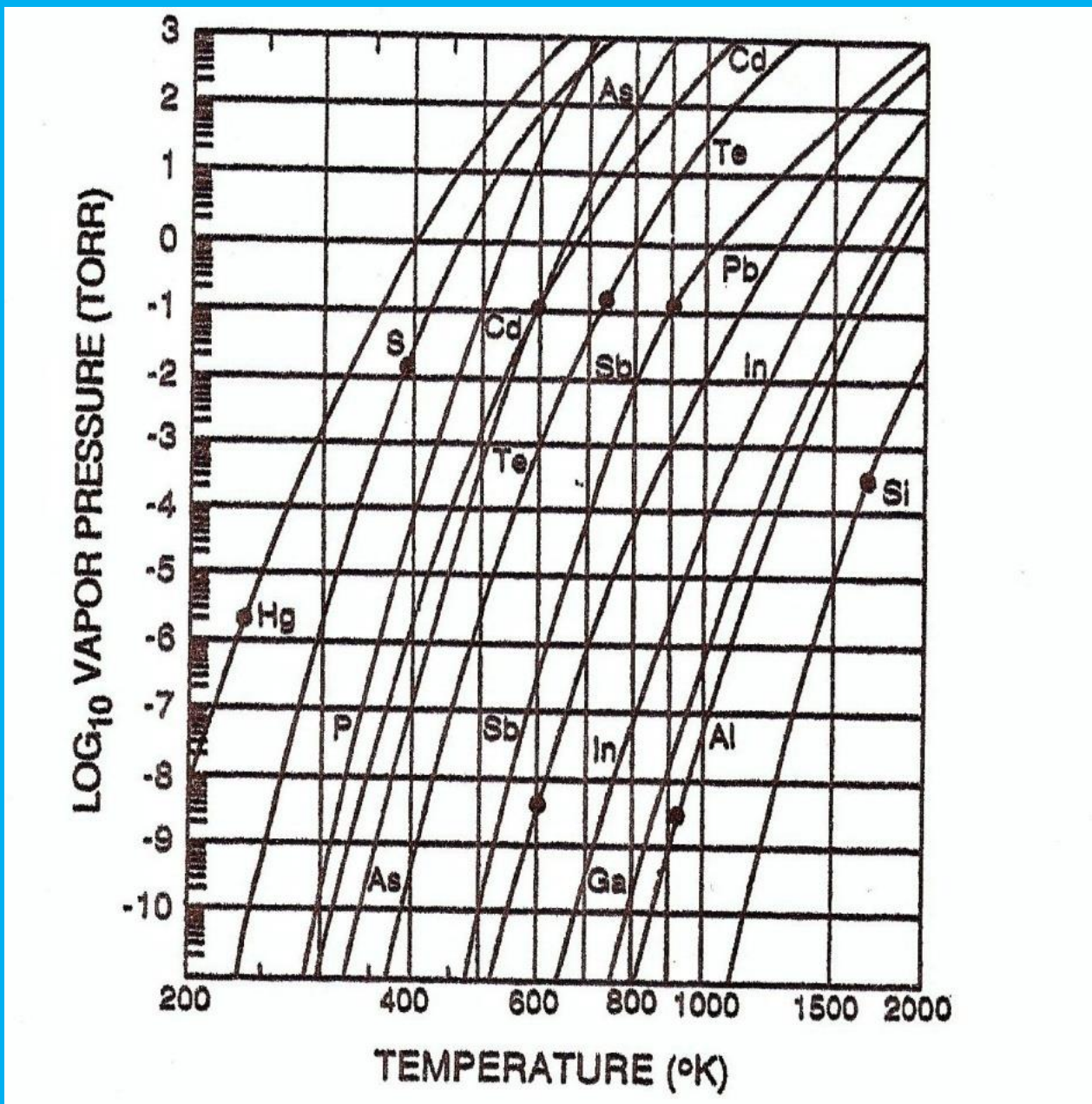
^g G. Thieme, *Vacuum*, 13, 55 (1963);

^h B. B. Dayton, *Trans. 6th Nat. Vac. Symp. Vac. Technol. (1959)*, Pergamon Press, New York, 1960, p.101.

Table 3-11 Gassing Rates at Room Temperature in Torr Liters/cm² sec

<i>Material</i>	<i>After a few hours of pumping</i>	<i>After a 24-hr bake</i>	<i>Bakeout temp., °C</i>	<i>Ref.</i>
Nylon 51	2×10^{-8} (51 hr)	4×10^{-11}	120	(59)
Araldite CT200 + HT901	5×10^{-8} (51 hr)	10^{-10}	100	(59)
Neoprene	10^{-6}			(57)
Viton A	7×10^{-8} (52 hr)	1.3×10^{-9}	200	(60)
Teflon	10^{-8} - 10^{-7}	Low	250	(57),(61)
Glass	10^{-9} - 10^{-8}	10^{-15} - 10^{-14}	400	(62)
Ceramic	10^{-9} - 10^{-8}	10^{-15} - 10^{-14}	400	(57),(62)
Metal	10^{-9} - 10^{-8} (10^{-10} after 50-100 hr)	10^{-15} - 10^{-14}	400	(62),(63)

Pressão de Vapor em função da Temperatura



Modelos de Fontes de Gases em um Sistema de Vácuo

Fonte de gás	Característica	Comentário
Volume	$P = P_o e^{-\frac{S}{V}t}$	Pressão cai exponencialmente dependendo de S e V
Vazamento Real	$P_{res} = \frac{Q_{vr}}{S}; Q_{vr} \approx C_{vr} P_{atm}$	Fluxo constante. Utilizar detector de vazamentos. Deve ser eliminado
Vazamento Virtual	$Q_{vv} = \frac{C_v}{S} P_o e^{-\frac{C_v}{V_c}t}$	$C_v \ll S b$ Queda da pressão depende de C_v e V_c . Evitar no projeto
Difusão	$Q(t) = c_o \frac{\sqrt{D}}{\sqrt{\pi t}}$	$Q(t)$ é proporcional a $\frac{1}{\sqrt{t}}$
Permeação	$Q = \frac{K(P_e^n - P_i^n)}{d}$	$N=1$ para não metais; $n=1/2$ para moléculas diatômicas em metais. Constante de permeação $K(T)$ é proporcional a $10^3/T$
Evaporação	$W = 0.058 P_v \sqrt{\frac{M}{T}} \frac{g}{cm^2 s}$ $Q = WA$ (g/s) $Q = \frac{\Delta N}{\Delta t} kT \frac{Torrl}{s}$	Crescimento de P_v em função da temperatura é exponencial e por isso mais rápido do que $\frac{1}{\sqrt{T}}$
Desorção Térmica (desgaseificação)	Primeira ordem: $\frac{dc}{dt} = c_o k_1 e^{-\frac{t}{\tau_{res}}}$ $\frac{1}{k_1} = \tau_{res} = \tau_o e^{\frac{E_d}{N k T}}$	Rápido $\tau_{res} = 10^{-12} s$
Temperatura (cozimento)	Segunda ordem: $\frac{dc}{dt} = \frac{-k_2 C_o^2}{(1 + C_o K_2 t)^2}$	Cai lentamente A molécula de H_2 se dissocia na adsorção e recombina na desorção
Superfícies Reais	$q_n = \frac{q}{t^\alpha}$ $0.7 \leq \alpha \leq 2$	Fórmula geral $q_n = q t^{-1}$ Adsorção química Adsorção física

4.6 PRESSURE LIMITS

75

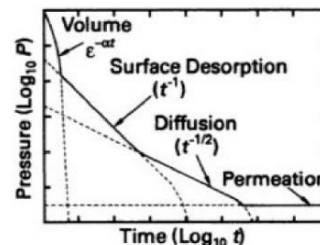


Fig. 4.10 Rate limiting steps during the pumping of a vacuum chamber.