

Introduction to Cryogenics

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Contents

- Introduction
- Cryogenic fluids
- Heat transfer & thermal insulation
- Cryogenic distribution & cooling schemes
- Refrigeration & liquefaction
- Cryogen storage & transport
- Thermometry

- **cryogenics**, that branch of physics which deals with the production of very low temperatures and their effects on matter

Oxford English Dictionary

2nd edition, Oxford University Press (1989)

- **cryogenics**, the science and technology of temperatures below 120 K

New International Dictionary of Refrigeration

3rd edition, IIF-IIR Paris (1975)

Characteristic temperatures of cryogens

Cryogen	Triple point [K]	Normal boiling point [K]	Critical point [K]
Methane	90.7	111.6	190.5
Oxygen	54.4	90.2	154.6
Argon	83.8	87.3	150.9
Nitrogen	63.1	77.3	126.2
Neon	24.6	27.1	44.4
Hydrogen	13.8	20.4	33.2
Helium	2.2 (*)	4.2	5.2

(*): λ Point

Cryogenic transport of natural gas: LNG



130 000 m³ LNG carrier
with double hull

Invar® tanks hold LNG at ~110 K



Densification, liquefaction & separation of gases

Ariane 5

25 t LH₂, 130 t LO₂



Space Shuttle

100 t LH₂, 600 t LO₂



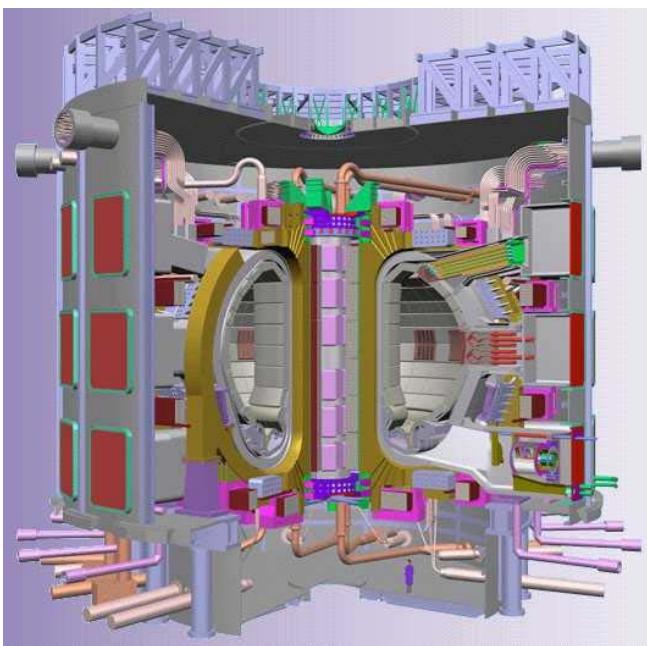
What are low temperatures?

- Entropy and temperature
 - the entropy of a thermodynamical system in a macrostate corresponding to a multiplicity Ω of microstates is
$$S = k_B \ln \Omega$$
 - adding reversibly heat dQ to the system results in a change of its entropy dS with a proportionality factor T
$$T = dQ/dS$$
 - ⇒ *high temperature: heating produces small entropy change*
 - ⇒ *low temperature: heating produces large entropy change*
- 1 K is equivalent to 10^{-4} eV or 10^{-23} J thermal energy
 - a temperature is « low » when $k_B T$ is small compared with the characteristic energy of the process considered
 - cryogenic temperatures reveal phenomena with low characteristic energy and enable their application

Characteristic temperatures of low-energy phenomena

Phenomenon	Temperature
Debye temperature of metals	few 100 K
High-temperature superconductors	~ 100 K
Low-temperature superconductors	~ 10 K
Intrinsic transport properties of metals	< 10 K
Cryopumping	few K
Cosmic microwave background	2.7 K
Superfluid helium 4	2.2 K
Bolometers for cosmic radiation	< 1 K
Low-density atomic Bose-Einstein condensates	~ μK

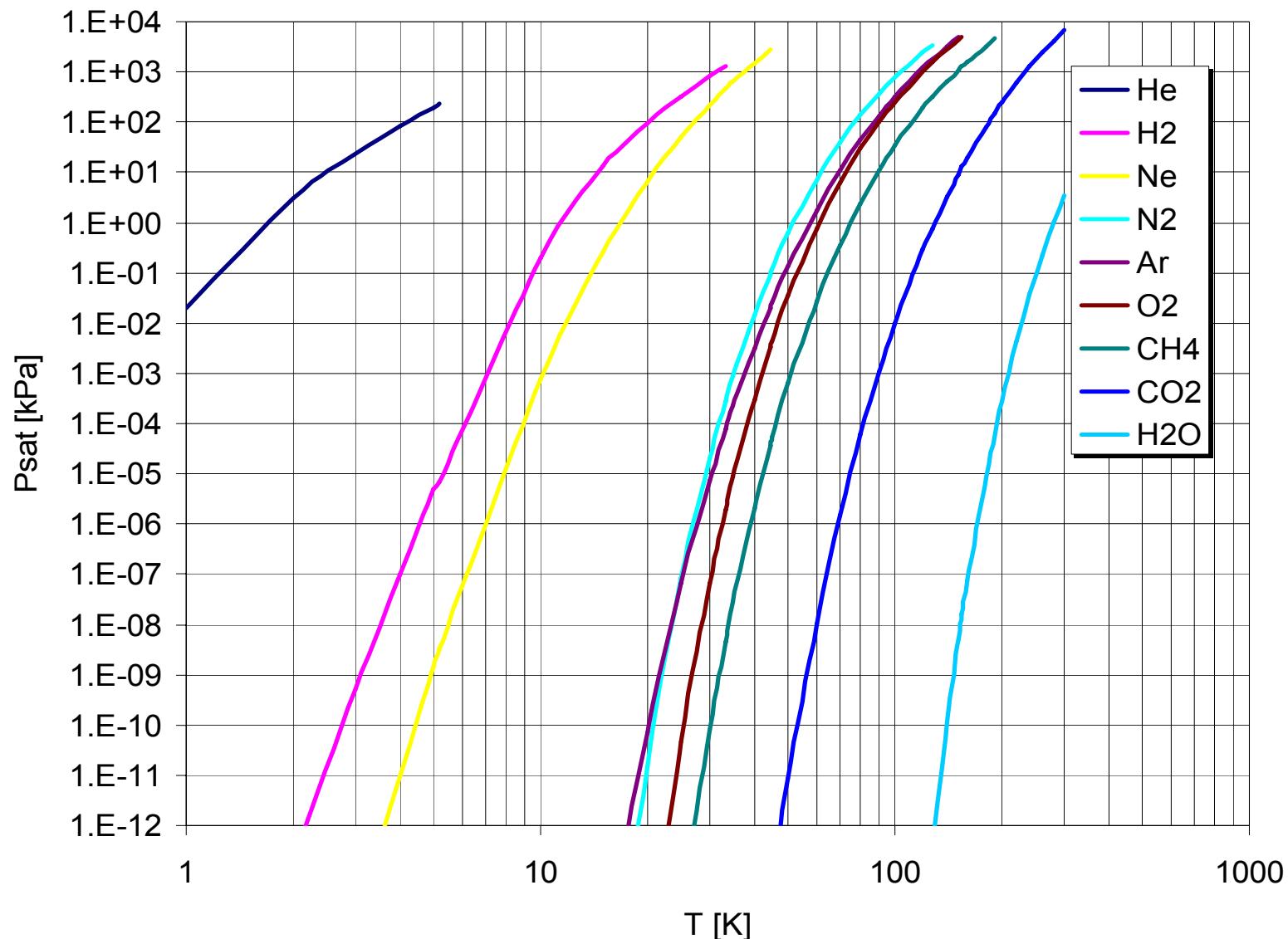
Cooling of superconducting devices



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Vapour pressure at cryogenic temperatures



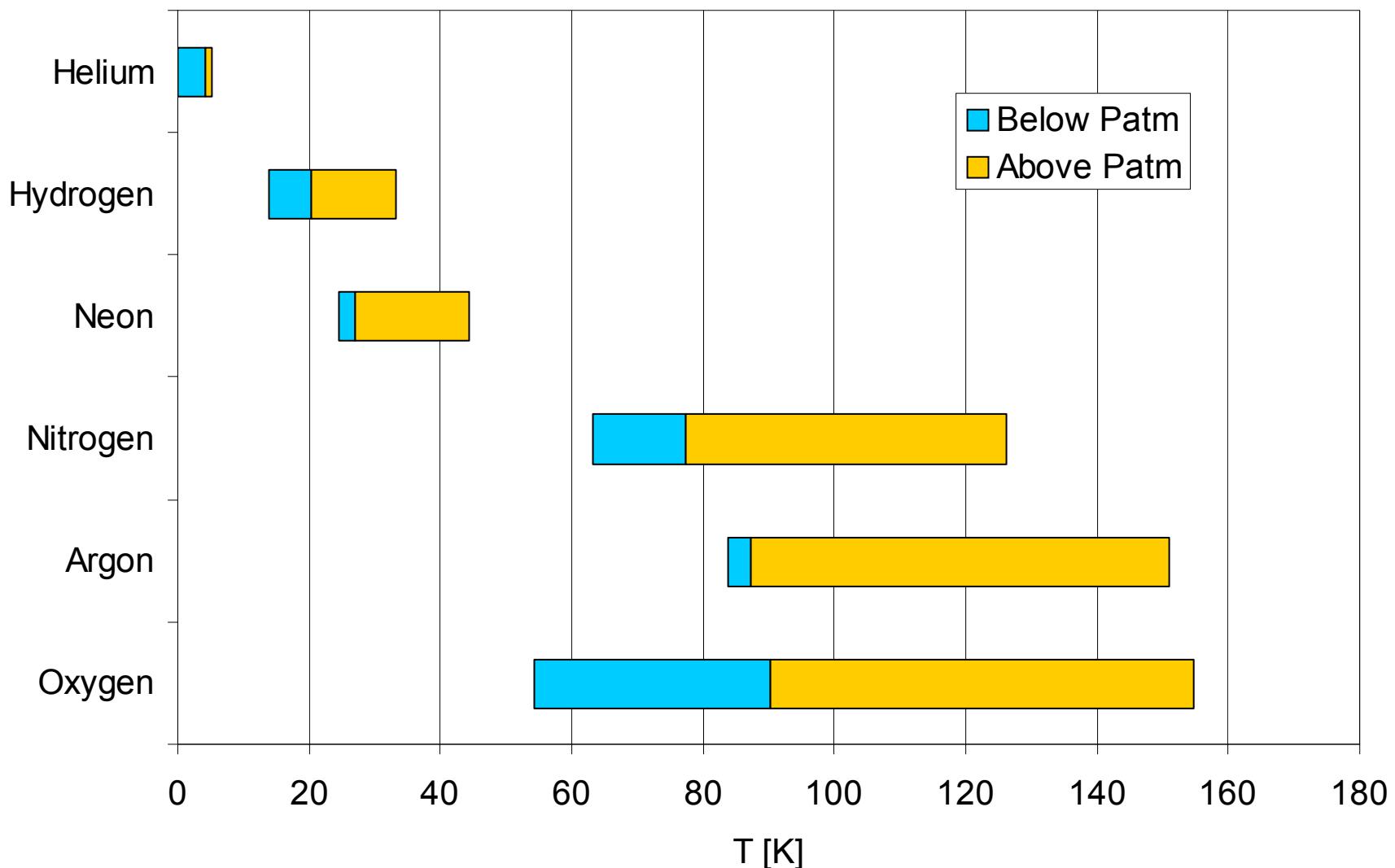
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Useful range of cryogens



Properties of cryogens compared to water

Property		He	N ₂	H ₂ O
Normal boiling point	[K]	4.2	77	373
Critical temperature	[K]	5.2	126	647
Critical pressure	[bar]	2.3	34	221
Liq./Vap. density (*)		7.4	175	1600
Heat of vaporization (*)	[J.g ⁻¹]	20.4	199	2260
Liquid viscosity (*)	[μPI]	3.3	152	278

(*) at normal boiling point

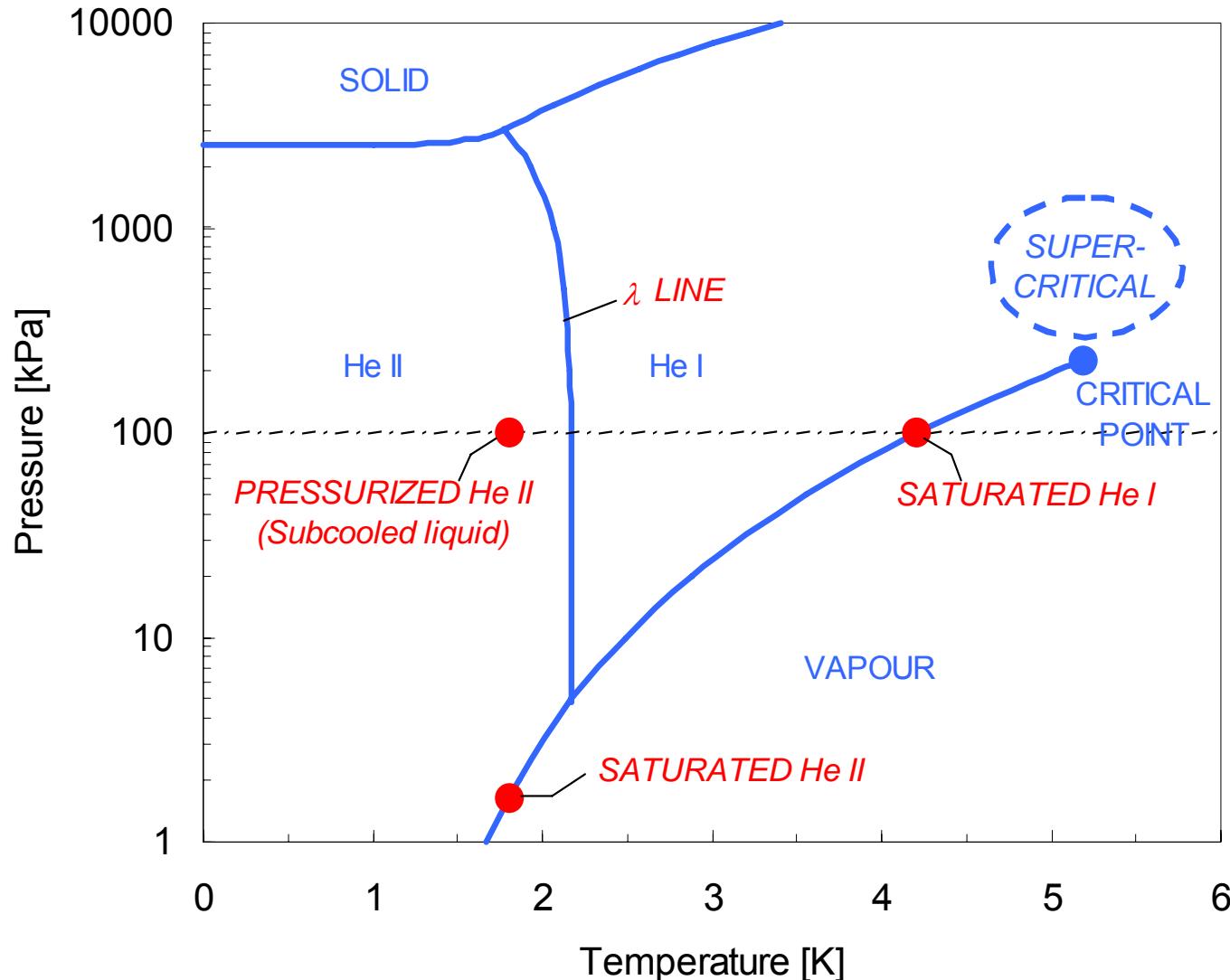
Vaporization of normal boiling cryogens under 1 W applied heat load

Cryogen	[mg.s ⁻¹]	[l.h ⁻¹] (liquid)	[l.min ⁻¹] (gas NTP)
Helium	48	1.38	16.4
Nitrogen	5	0.02	0.24

Amount of cryogens required to cool down 1 kg iron

Using	Latent heat only	Latent heat and enthalpy of gas
LHe from 290 to 4.2 K	29.5 litre	0.75 liter
LHe from 77 to 4.2 K	1.46 litre	0.12 litre
LN2 from 290 to 77 K	0.45 litre	0.29 litre

Phase diagram of helium



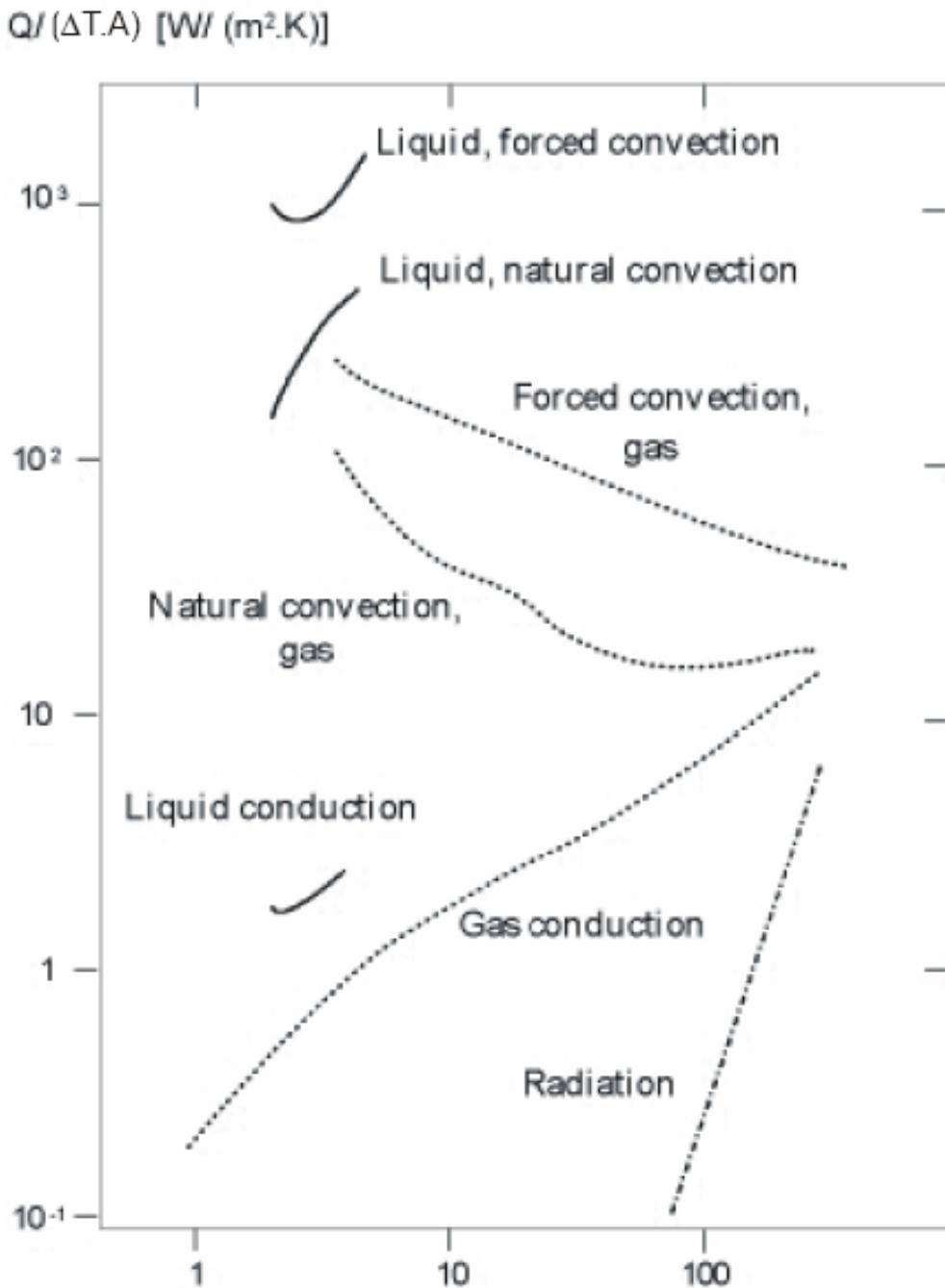
Helium as a cooling fluid

Phase domain	Advantages	Drawbacks
Saturated He I	Fixed temperature High heat transfer	Two-phase flow Boiling crisis
Supercritical	Monophase Negative J-T effect	Non-isothermal Density wave instability
He II	Low temperature High conductivity Low viscosity	Second-law cost Subatmospheric

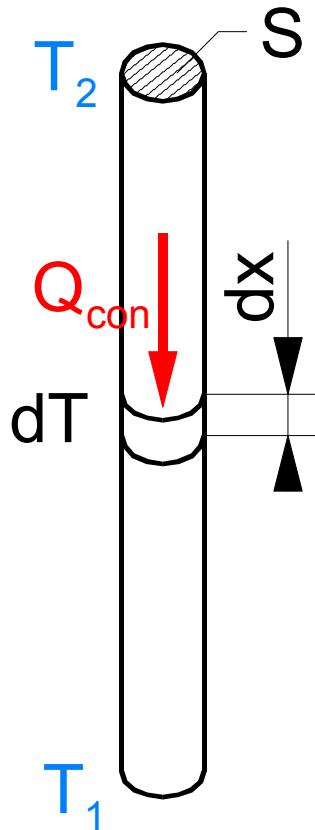
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Typical heat transfer coefficients at cryogenic temperatures



Heat conduction in solids



Fourier's law:

$$Q_{\text{con}} = k(T) \cdot S \cdot \frac{dT}{dx}$$

k(T): thermal conductivity [W/m.K]

Integral form:

$$Q_{\text{con}} = \frac{S}{L} \cdot \int_{T_1}^{T_2} k(T) \cdot dT$$

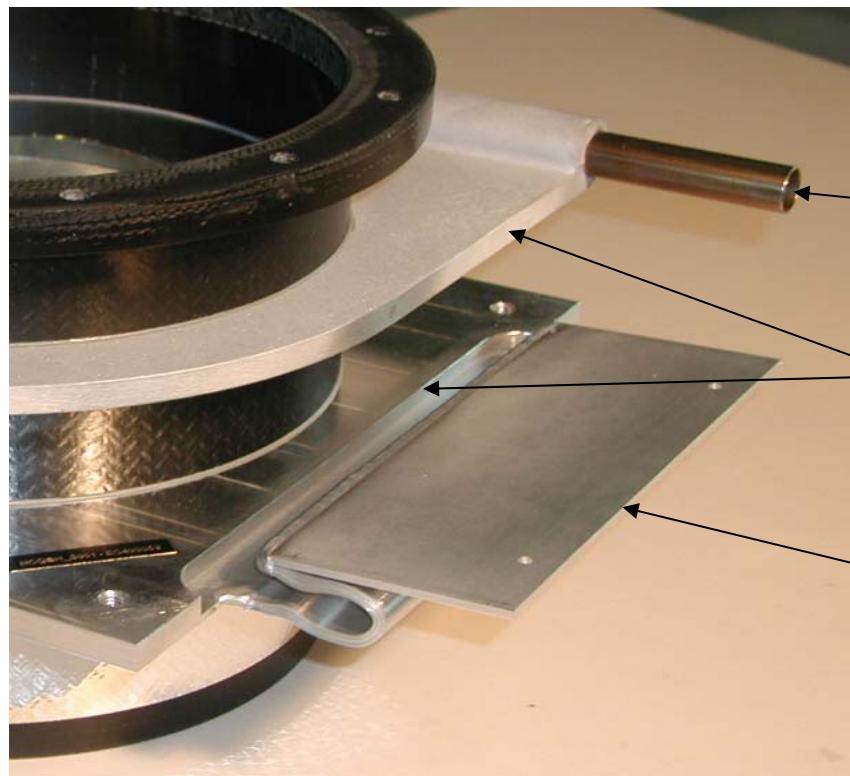
$\int k(T) \cdot dT$: thermal conductivity integral [W/m]

Thermal conductivity integrals for standard construction materials are tabulated

Thermal conductivity integrals of selected materials [W/m]

From vanishingly low temperature up to	20 K	80 K	290 K
OFHC copper	11000	60600	152000
DHP copper	395	5890	46100
1100 aluminium	2740	23300	72100
2024 aluminium alloy	160	2420	22900
AISI 304 stainless steel	16.3	349	3060
G-10 glass-epoxy composite	2	18	153

Non-metallic composite support post with heat intercepts



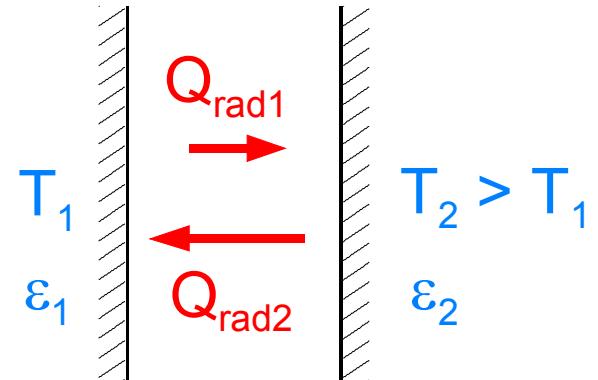
5 K cooling line (SC He)

Aluminium intercept plates
glued to G-10 column

Aluminium strips to thermal
shield at 50-75 K

Thermal radiation

- Wien's law
 - Maximum of black body power spectrum
 $\lambda_{max} T = 2898 \text{ [\mu m.K]}$
- Stefan-Boltzmann's law
 - Black body
 - "Gray"body
 - "Gray" surfaces at T_1 and T_2



$$Q_{rad} = \sigma A T^4$$

$$\sigma = 5.67 \times 10^{-8} \text{ W/m}^2\text{.K}^4$$

(Stefan Boltzmann's constant)

$$Q_{rad} = \varepsilon \sigma A T^4$$

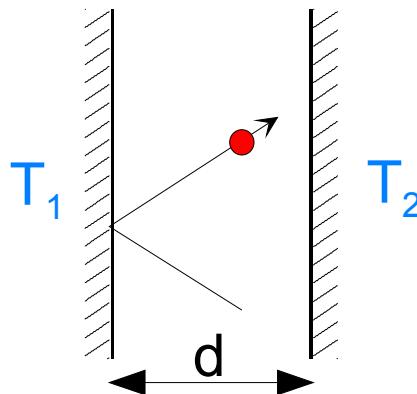
ε emissivity of surface

$$Q_{rad} = E \sigma A (T_1^4 - T_2^4)$$

E function of $\varepsilon_1, \varepsilon_2$, geometry

Emissivity of technical materials at low temperatures

	Radiation from 290 K Surface at 77 K	Radiation from 77 K Surface at 4.2 K
Stainless steel, as found	0.34	0.12
Stainless steel, mech. polished	0.12	0.07
Stainless steel, electropolished	0.10	0.07
Stainless steel + Al foil	0.05	0.01
Aluminium, as found	0.12	0.07
Aluminium, mech. polished	0.10	0.06
Aluminium, electropolished	0.08	0.04
Copper, as found	0.12	0.06
Copper, mech. Polished	0.06	0.02



Residual gas conduction

$\lambda_{molecule}$: mean free path of gas molecules

- Viscous regime

- At high gas pressure $\lambda_{molecule} \ll d$
- Classical conduction $Q_{res} = k(T) A dT/dx$
- Thermal conductivity $k(T)$ independant of pressure

- Molecular regime

- At low gas pressure $\lambda_{molecule} \gg d$
- Kennard's law $Q_{res} = A \alpha(T) \Omega P (T_2 - T_1)$
- Conduction heat transfer proportional to pressure, independant of spacing between surfaces
 Ω depends on gas species
- Accommodation coefficient $\alpha(T)$ depends on gas species, T_1 , T_2 , and geometry of facing surfaces

Multi-layer insulation (MLI)



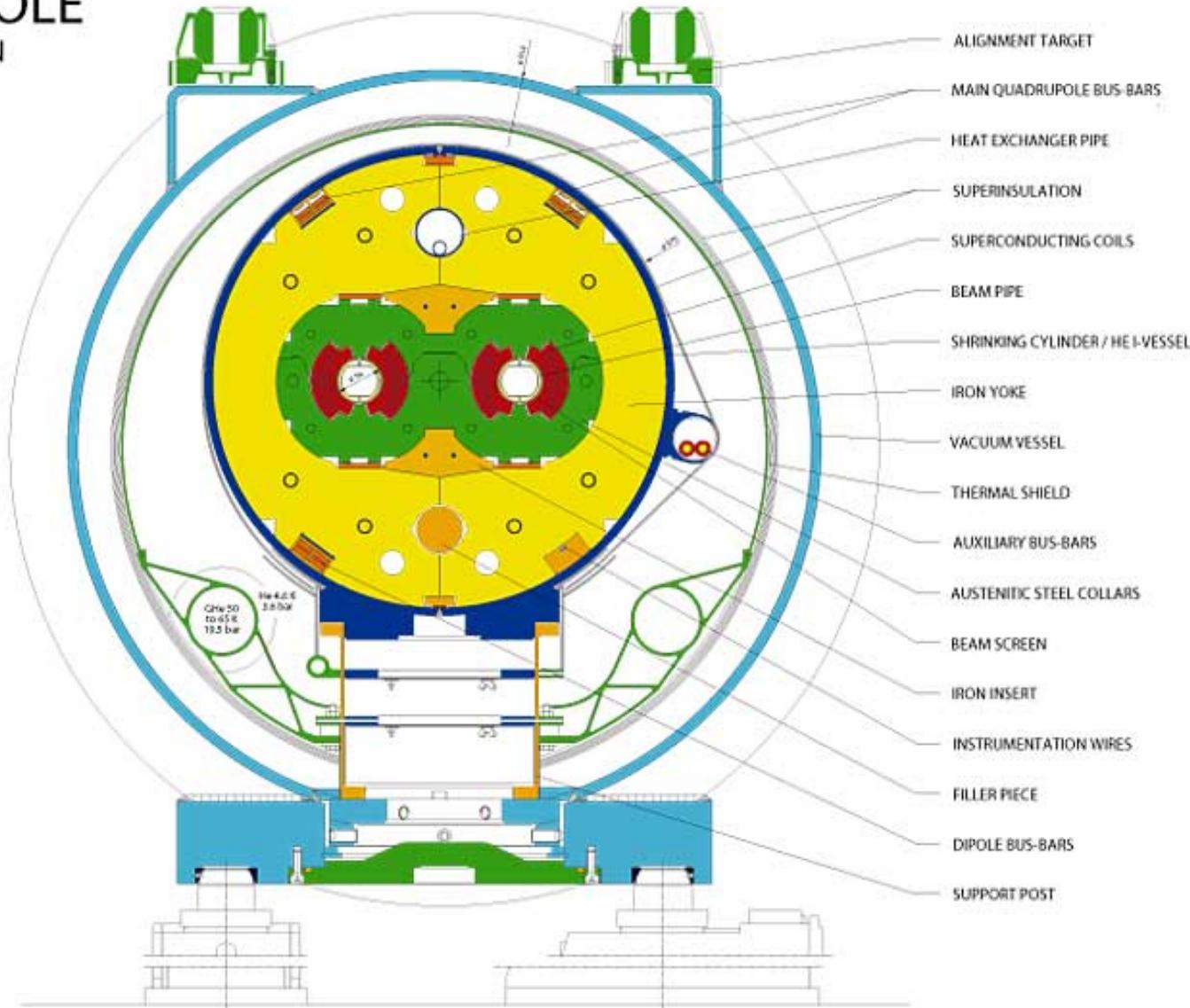
- Complex system involving three heat transfer processes
 - $Q_{MLI} = Q_{rad} + Q_{sol} + Q_{res}$
 - With n reflective layers of equal emissivity, $Q_{rad} \sim 1/(n+1)$
 - Due to parasitic contacts between layers, Q_{sol} increases with layer density
 - Q_{res} due to residual gas trapped between layers, scales as $1/n$ in molecular regime
 - Non-linear behaviour requires layer-to-layer modeling
- In practice
 - Typical data available from (abundant) literature
 - Measure performance on test samples

Typical heat fluxes at vanishingly low temperature between flat plates [W/m²]

Black-body radiation from 290 K	401
Black-body radiation from 80 K	2.3
Gas conduction (100 mPa He) from 290 K	19
Gas conduction (1 mPa He) from 290 K	0.19
Gas conduction (100 mPa He) from 80 K	6.8
Gas conduction (1 mPa He) from 80 K	0.07
MLI (30 layers) from 290 K, pressure below 1 mPa	1-1.5
MLI (10 layers) from 80 K, pressure below 1 mPa	0.05
MLI (10 layers) from 80 K, pressure 100 mPa	1-2

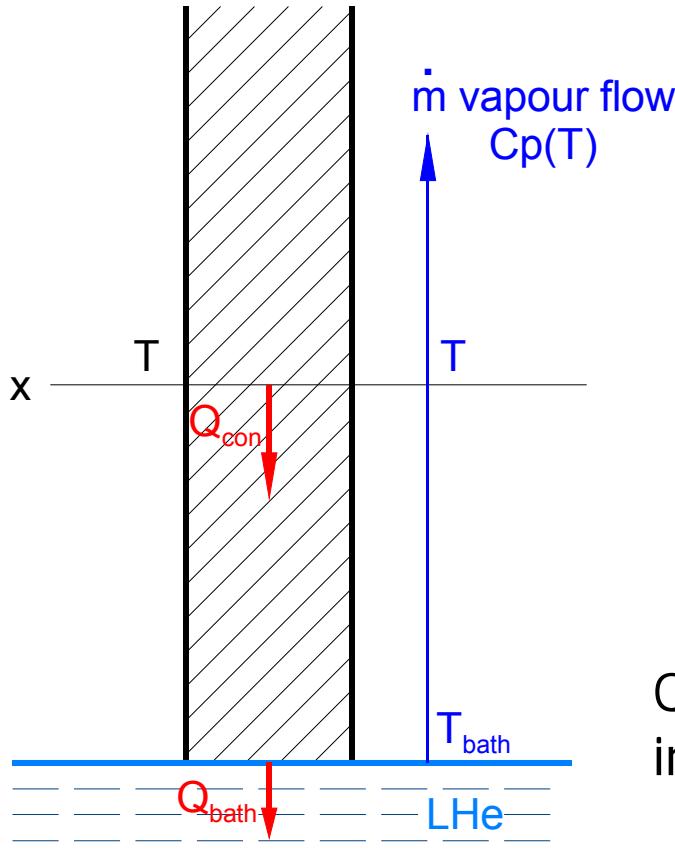
Cross-section of LHC dipole cryostat

LHC DIPOLE
CROSS SECTION



Vapour cooling of necks and supports with perfect heat exchange

Cross-section A



Assuming perfect heat exchange between solid and gas, i.e. $T_{\text{sol}}(x)=T_{\text{gas}}(x)=T(x)$:

$$Q_{\text{con}} = Q_{\text{bath}} + \dot{m} \cdot C_p(T) \cdot (T - T_{\text{bath}})$$

$$k(T) \cdot A \cdot \frac{dT}{dx} = Q_{\text{bath}} + \dot{m} \cdot C_p(T) \cdot (T - T_{\text{bath}})$$

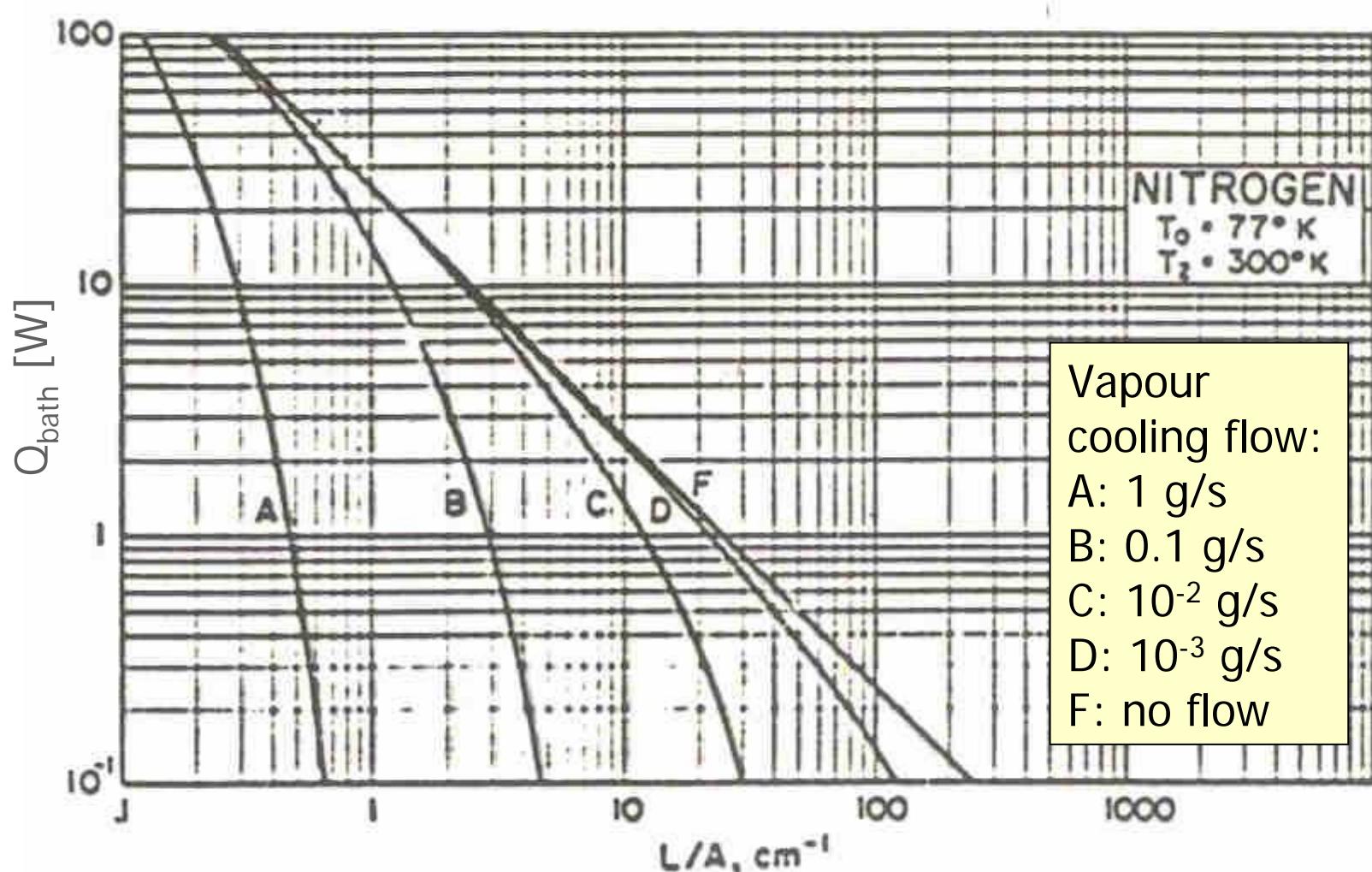
$C_p(T)$: Specific heat of vapour

$k(T)$: Thermal conductivity of the support

Q_{bath} can then be calculated by numerical integration for :

- different cryogens,
- different values of aspect ratio L/A
- different values of vapour flow

Heat reaching the cold end of a stainless steel neck



Vapour cooling of necks and supports with perfect heat exchange in self-sustained mode

A particular case of gas cooling is the self-sustained mode, i.e. He vapour flow is generated only by the residual heat Q_{bath} reaching the bath. Then:

$$Q_{\text{bath}} = L_v \cdot \dot{m} \quad (\text{Lv: latent heat of vaporization})$$

Given the general equation

$$k(T) \cdot A \cdot \frac{dT}{dx} = Q_{\text{bath}} + \dot{m} \cdot Cp(T) \cdot (T - T_{\text{bath}})$$

And after integration, we finally have:

$$Q_{\text{bath}} = \frac{A}{L} \cdot \int_{T_{\text{bath}}}^{T_{\text{ambient}}} \frac{K(T)}{1 + (T - T_{\text{bath}}) \cdot \frac{Cp(T)}{Lv}} \cdot dT$$

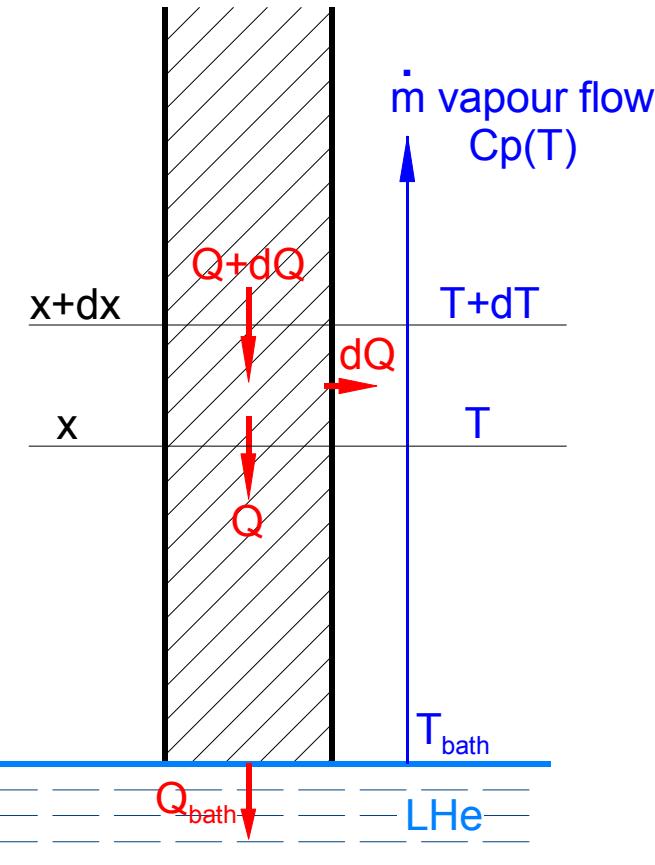
Attenuation factor w.r.
to pure conduction

Reduction of heat conduction by self-sustained helium vapour cooling

Effective thermal conductivity integral from 4 to 300 K	Purely conductive regime [W.cm ⁻¹]	Self-sustained vapour-cooling [W.cm ⁻¹]
ETP copper	1620	128
OFHC copper	1520	110
Aluminium 1100	728	39.9
Nickel 99% pure	213	8.65
Constantan	51.6	1.94
AISI 300 stainless steel	30.6	0.92

Vapour cooling of necks and supports with imperfect heat exchange

Cross-section A



$$dQ = f \cdot \dot{m} \cdot C_p(T) \cdot dT$$

With f , the efficiency of the heat transfer

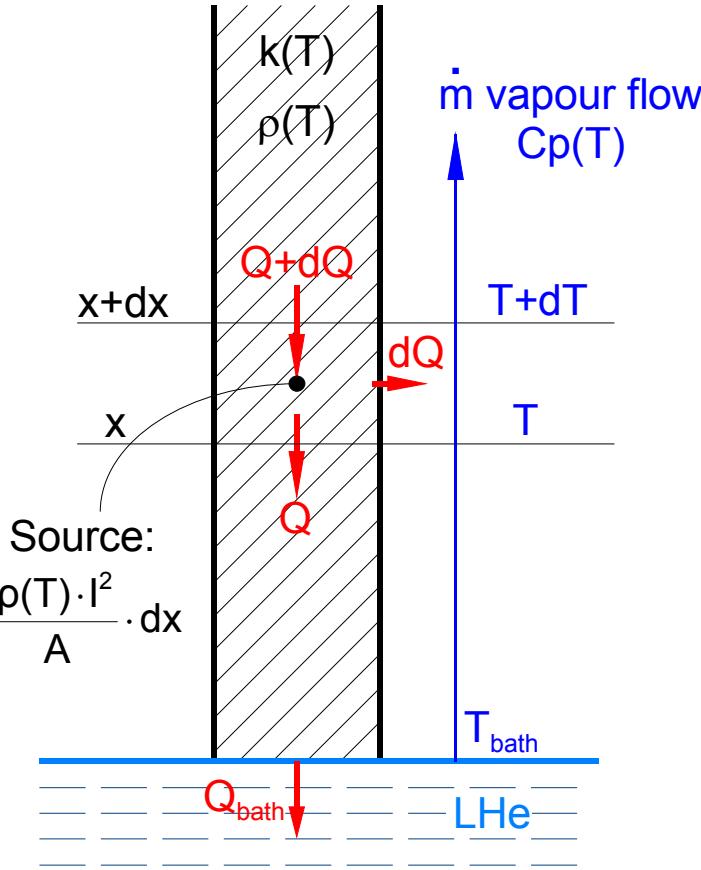
In steady state, the heat balance equation becomes:

$$\frac{d}{dx} \left[k(T) \cdot A \cdot \frac{dT}{dx} \right] = f \cdot \dot{m} \cdot C_p(T) \cdot \frac{dT}{dx}$$

→ Numerical integration for solving this equation

Vapor-cooled current leads

Cross-section A
Current I



$\rho(T)$: electrical resistivity

$$dQ = f \cdot \dot{m} \cdot Cp(T) \cdot dT$$

In steady-state, heat balance equation:

$$\frac{d}{dx} \left[k(T) \cdot A \cdot \frac{dT}{dx} \right] - f \cdot \dot{m} \cdot Cp(T) \cdot \frac{dT}{dx} + \frac{\rho(T) \cdot I^2}{A} = 0$$

Solid
conduction

Vapour
cooling

Joule
heating

Assuming the material of the lead follows the Wiedemann-Franz-Lorenz (WFL) law:

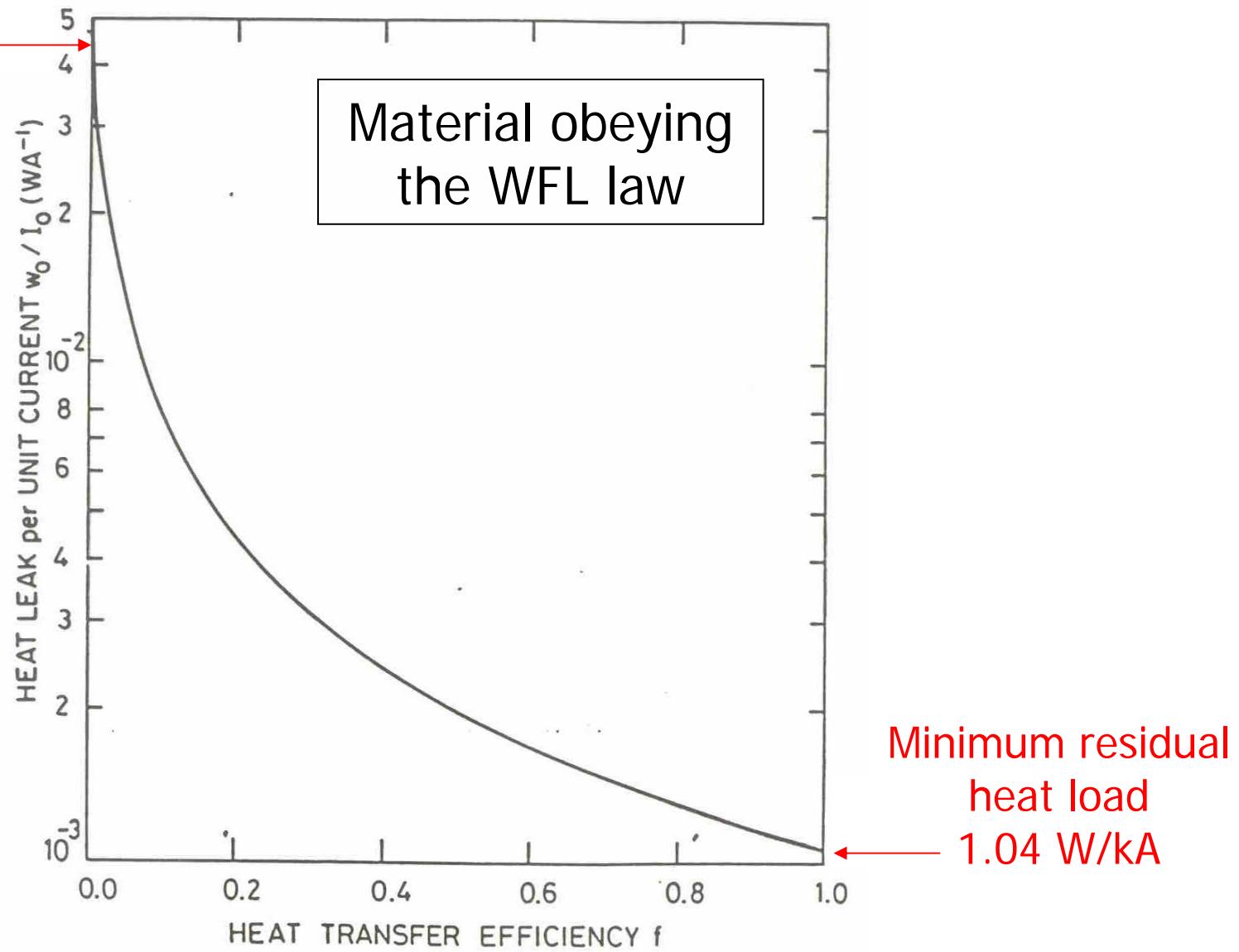
$$k(T) \cdot \rho(T) = L_0 \cdot T$$

L_0 : Lorenz number ($2.45 \cdot 10^{-8} \text{ W} \cdot \Omega \cdot \text{K}^{-2}$)

→ Then numerical integration

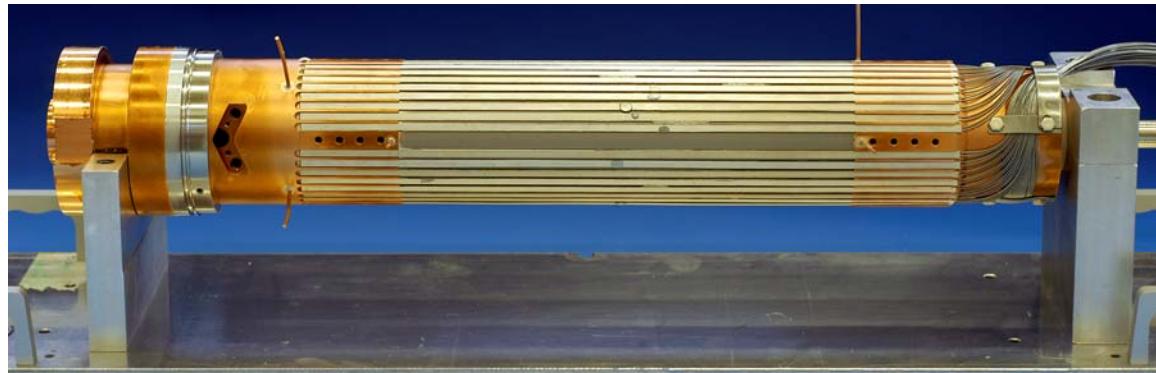
Heat load of optimized current lead

Uncooled
47 W/kA



Beating the WFL law: HTS current leads

- The WFL law essentially states that good electrical conductors are also good thermal conductors
- Current leads need good electrical conductors with low thermal conductivity
- Superconductors are bad thermal conductors with zero resistivity
- Build current lead with superconductor up to temperature as high as possible, i.e. HTS



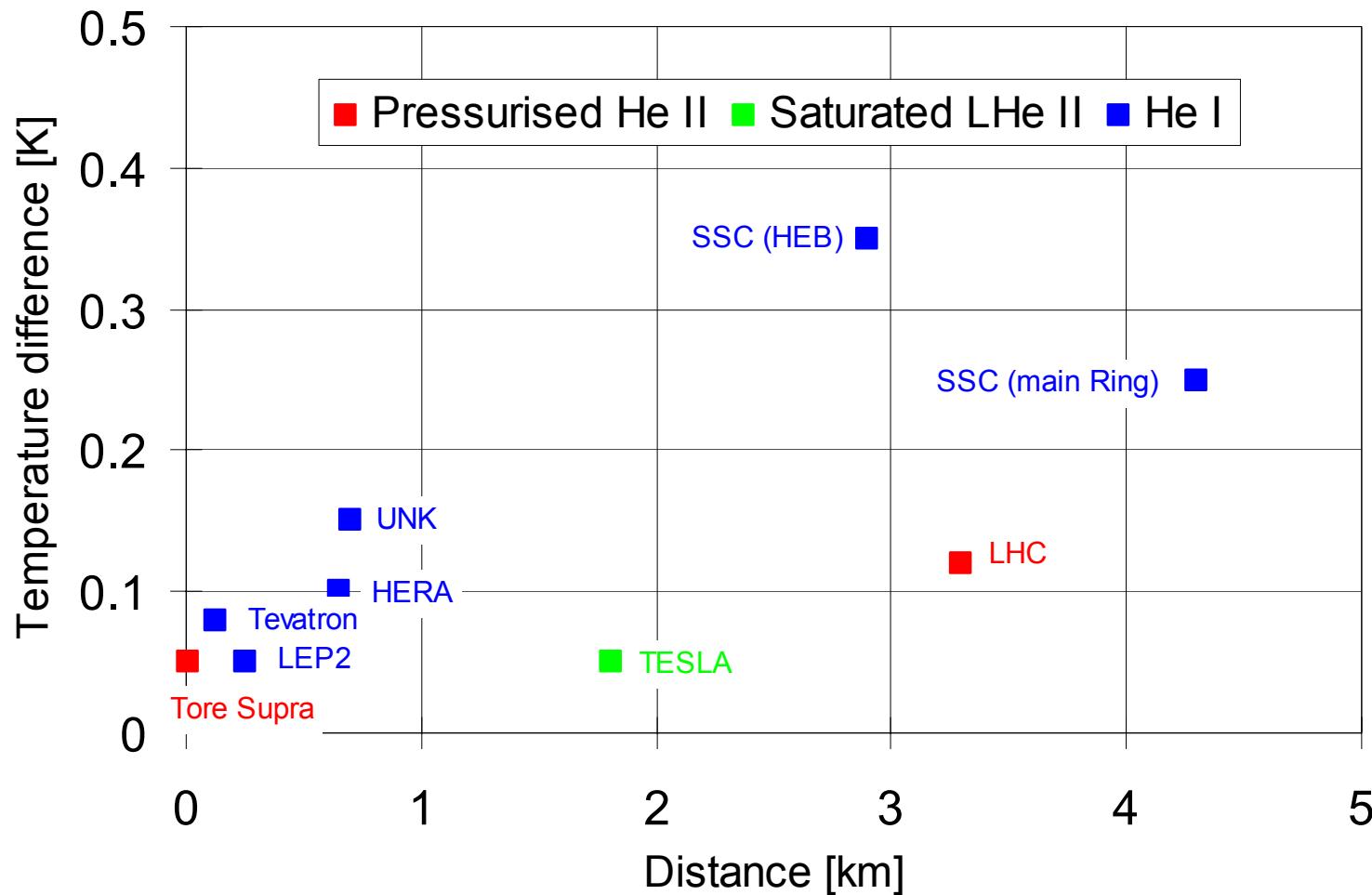
HTS vs. normal conducting current leads

Type	Resistive	HTS (4 to 50 K) Resistive (above)
Heat into LHe [W/kA]	1.1	0.1
Total exergy consumption [W/kA]	430	150
Electrical power from grid [W/kA]	1430	500

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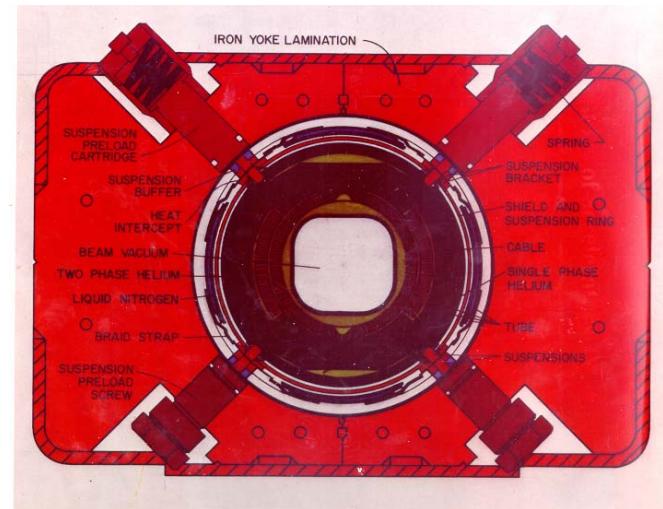
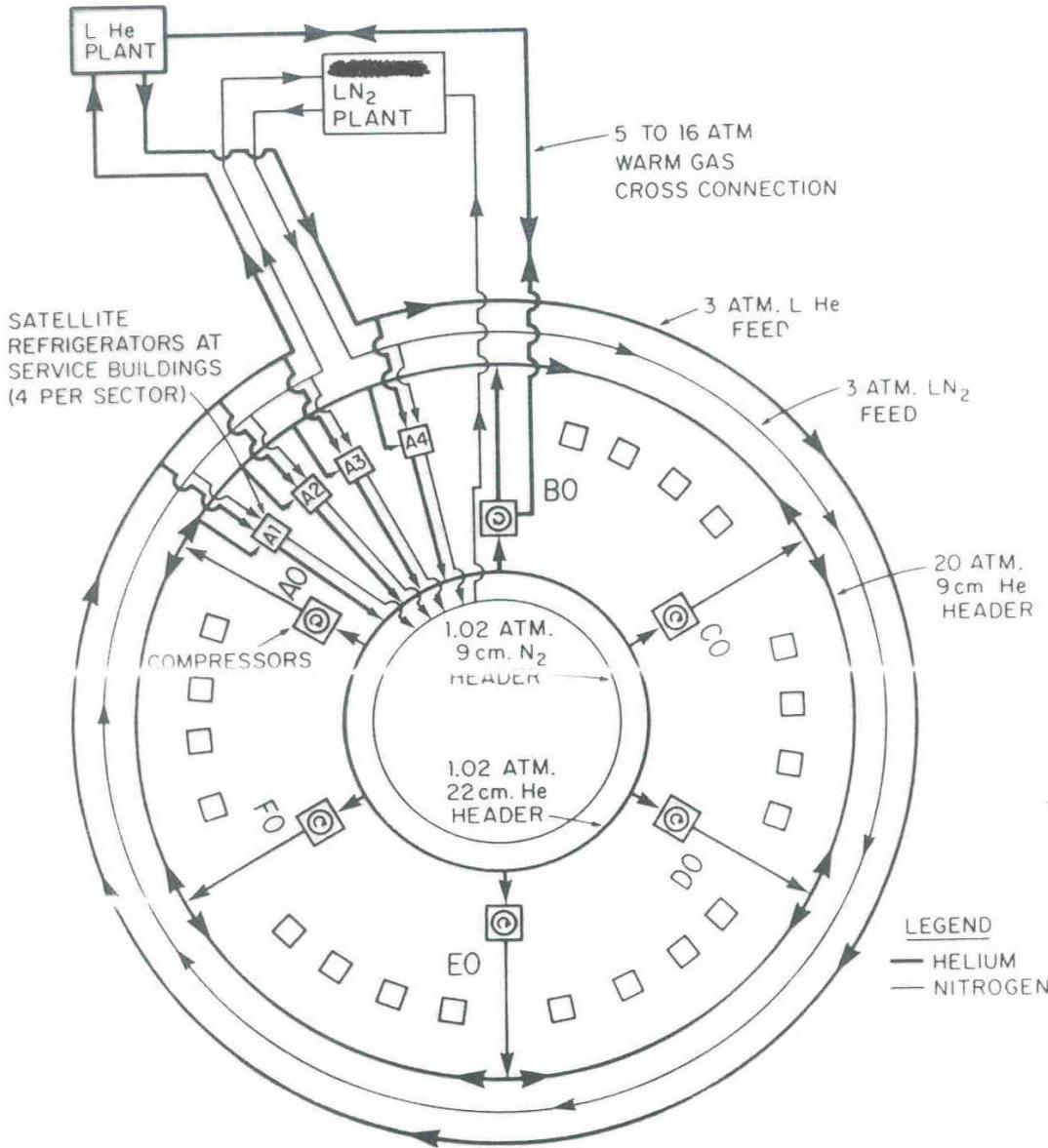
Transport of refrigeration in large distributed cryogenic systems



Cryogenic distribution scheme: design issues

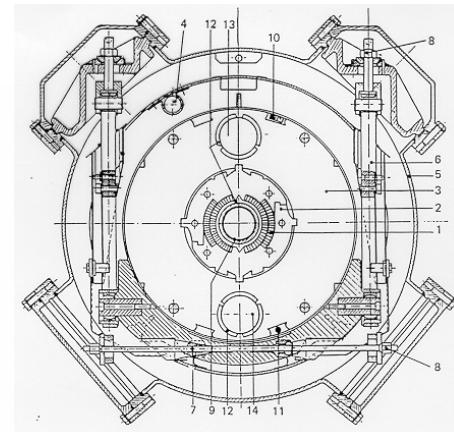
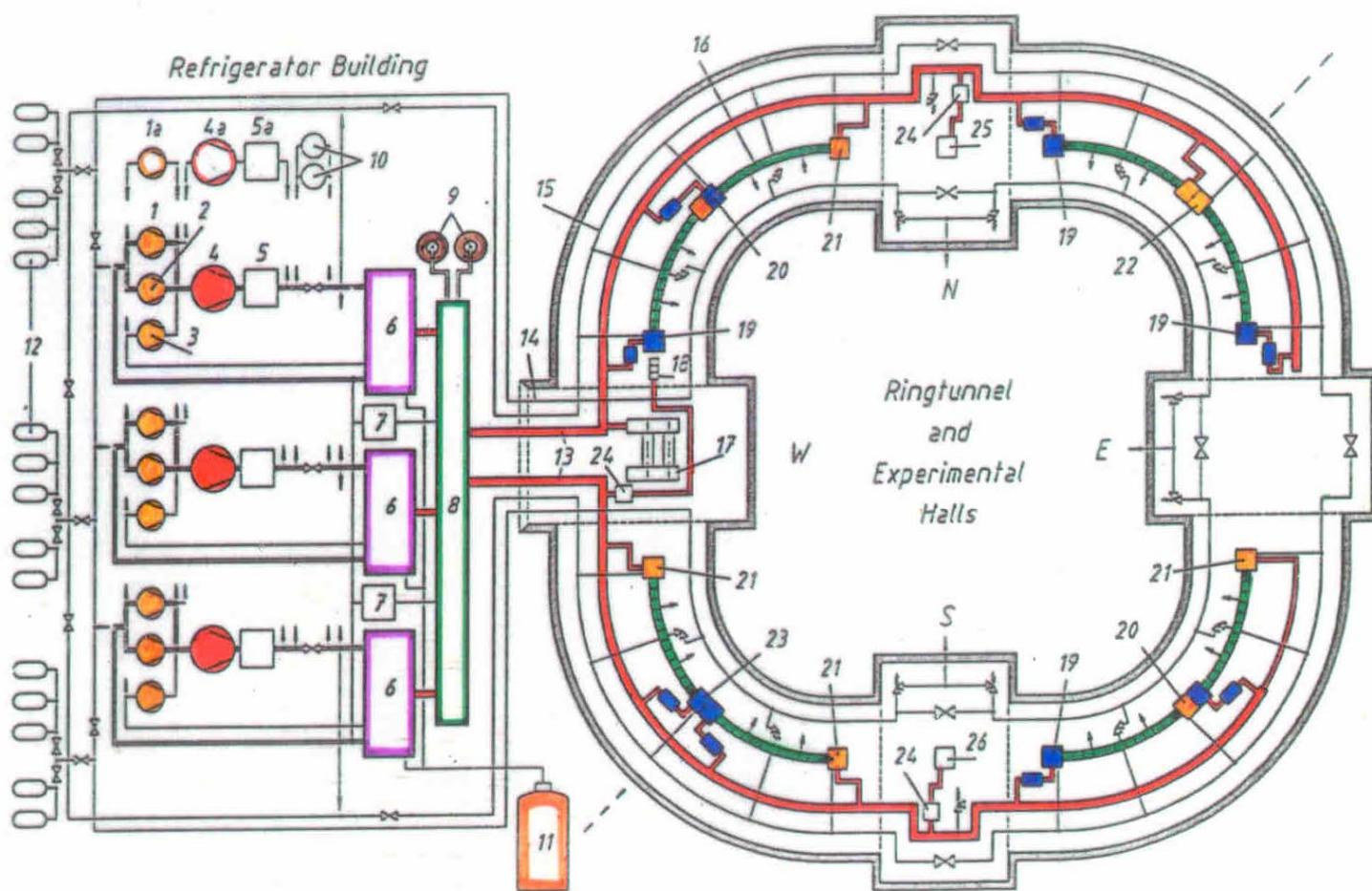
- Monophase vs. two-phase
 - temperature control
 - hydrostatic head & flow instabilities
- Pumps vs. no pumps
 - efficiency & cost
 - reliability & safety
- LN₂
 - cooldown and/or normal operation
 - capital & operating costs of additional fluid
 - safety in underground areas (ODH)
- Lumped vs. distributed cryoplants
- Separate cryoline vs. integrated piping
- Number of active components (valves, actuators)
- Redundancy of configuration

Tevatron distribution scheme



Central helium liquefier,
separate ring cryoline
and satellite
refrigerators

HERA distribution scheme



Central
cryoplant and
separate ring
cryoline

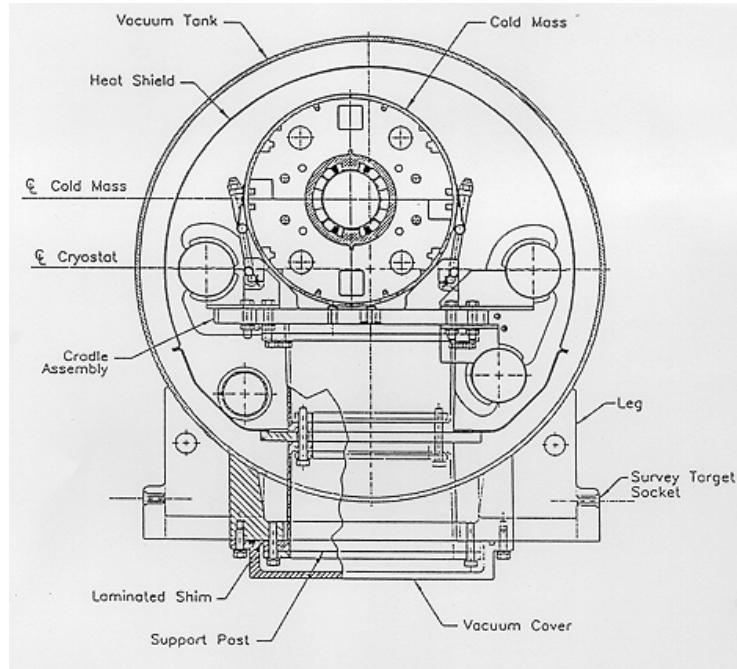
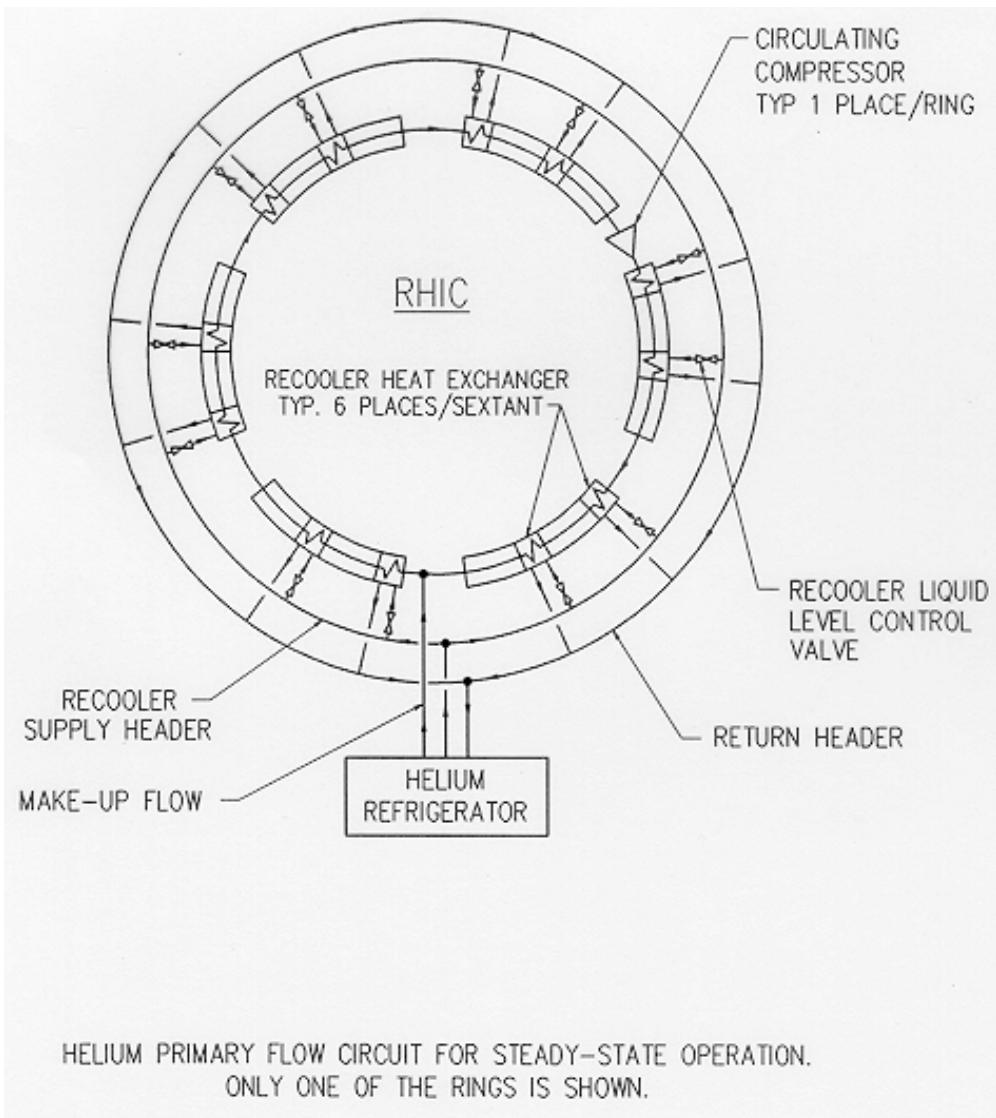
Refrigeration 4.3 K
Refrigeration 40/80 K
Current lead flow

6775 W
20000 W
 20.5×10^{-3} kg/s

total mass flow
Primary power
Specif. power consumption

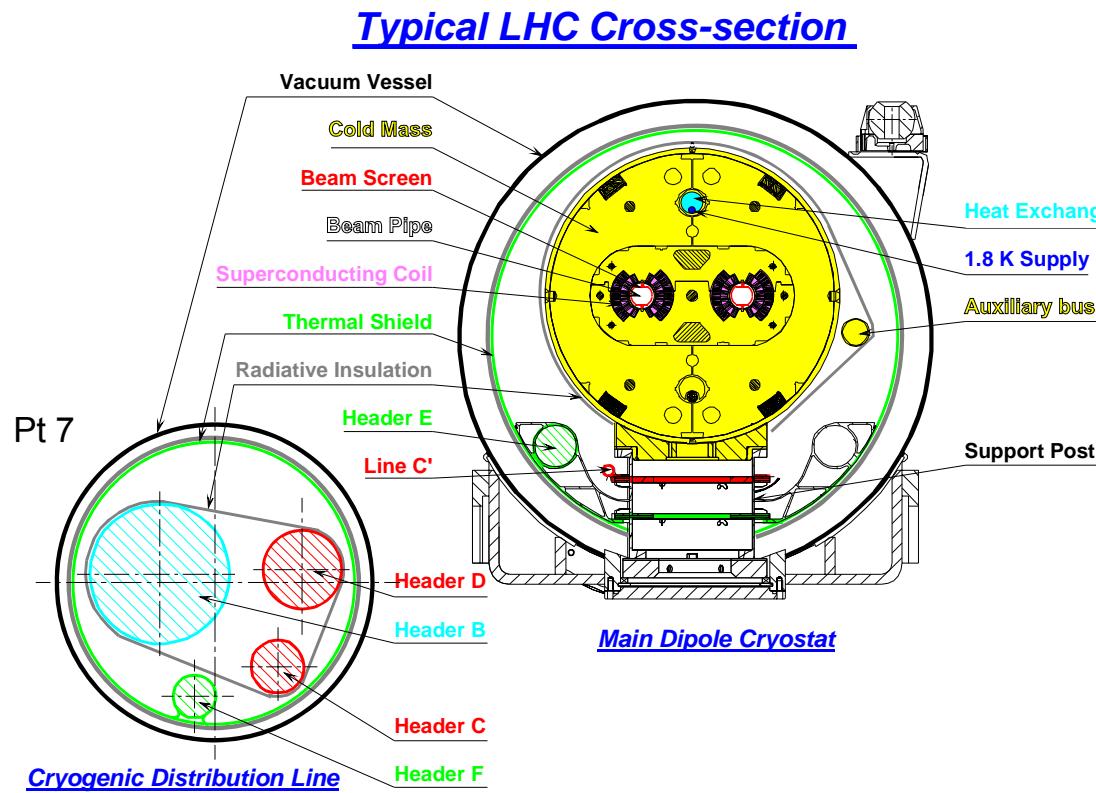
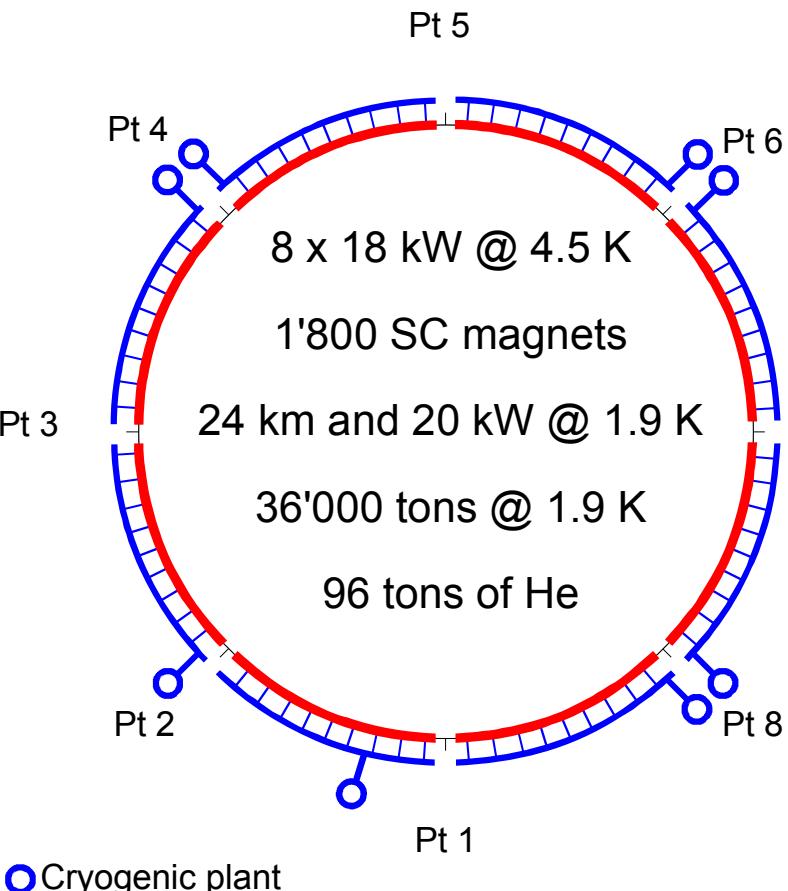
0.871 kg/s
2845 kW
281 W (300 K)/W (4.3 K)

RHIC distribution scheme



Central cryoplant and piping integrated in magnet cryostat

LHC distribution scheme

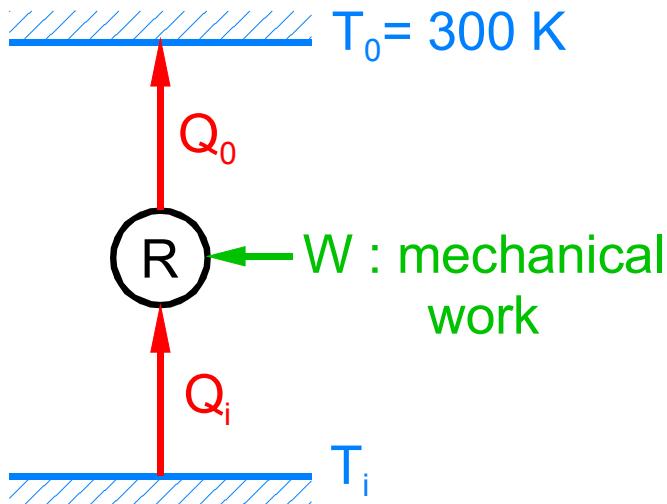


Cryoplants at
five points,
separate ring
cryoline

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Thermodynamics of cryogenic refrigeration



First principle [Joule]

$$Q_0 = Q_i + W$$

Second principle [Clausius]

$$\frac{Q_0}{T_0} \geq \frac{Q_i}{T_i}$$

(= for reversible process)

Hence, $W \geq T_0 \cdot \frac{Q_i}{T_i} - Q_i$

which can be written in three different ways:

① $W \geq T_0 \cdot \Delta S_i - Q_i$ introducing entropy S as

$$\Delta S_i = \frac{Q_i}{T_i}$$

② $W \geq Q_i \cdot \left(\frac{T_0}{T_i} - 1 \right)$ Carnot factor

③ $W \geq \Delta E_i$ introducing exergy E as

$$\Delta E_i = Q_i \cdot \left(\frac{T_0}{T_i} - 1 \right)$$

Minimum refrigeration work

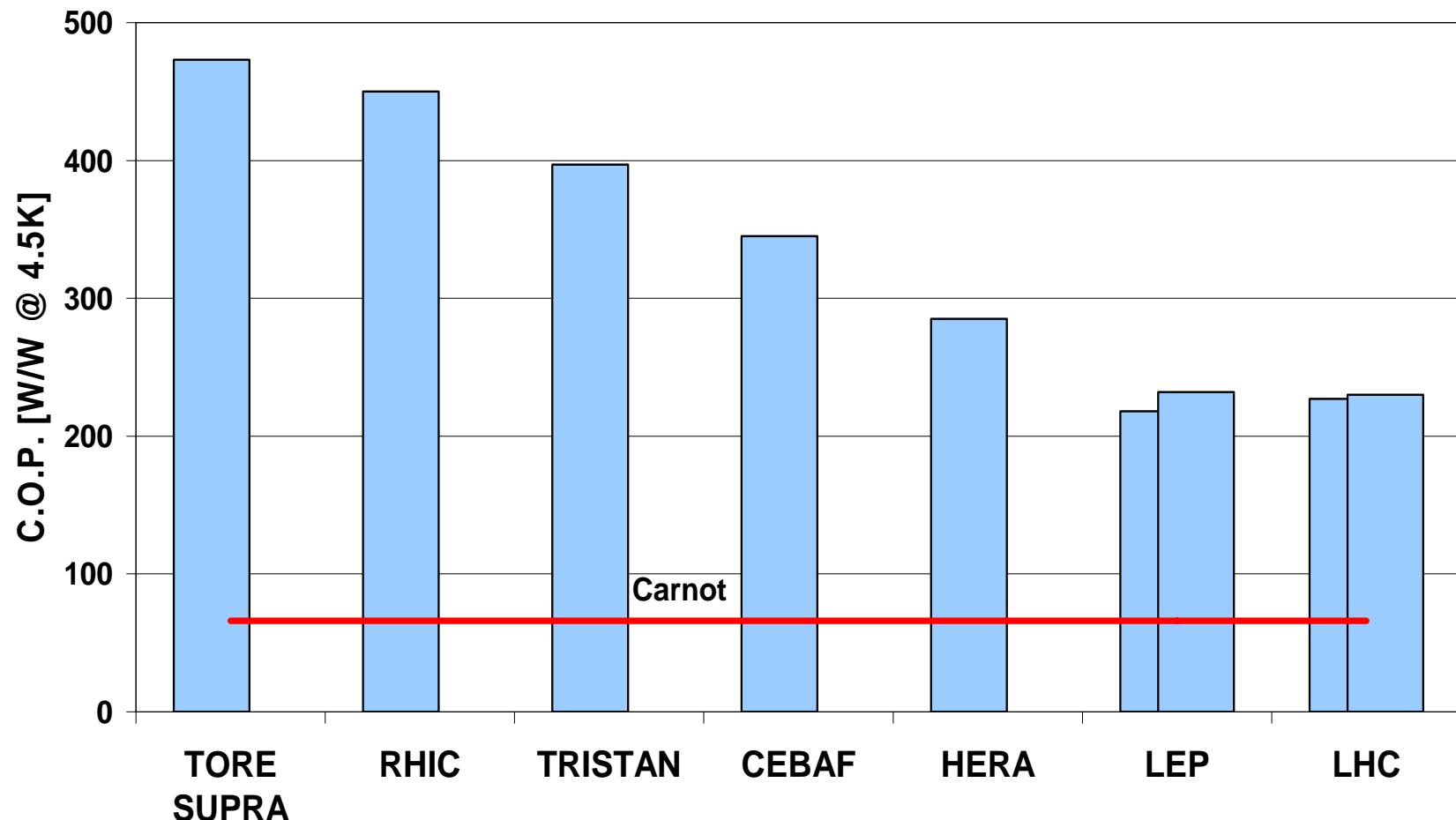
Consider the extraction of 1 W at 4.5 K, rejected at 300 K
The minimum refrigeration work (equation 2) is:

$$W_{\min} = Q_i \cdot \left(\frac{T_0}{T_i} - 1 \right) = 1 \cdot \left(\frac{300}{4.5} - 1 \right) = 65.7 \text{ W}$$

In practice, the most efficient helium refrigerators have an efficiency of about 30% w.r. to the Carnot limit.

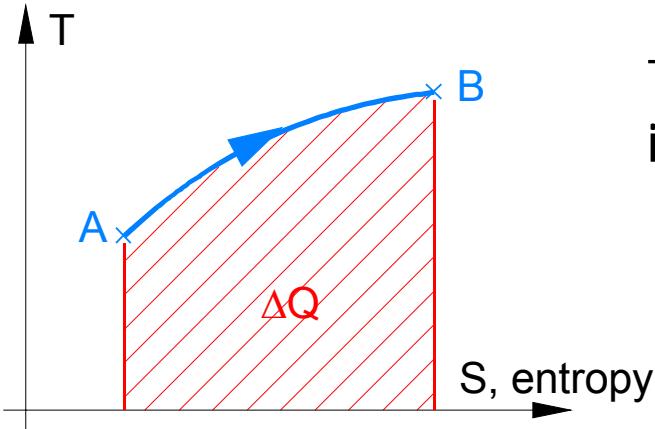
$$\Rightarrow W_{\text{real}} = \frac{W_{\min}}{\eta} = \frac{65.7}{0.3} = 220 \text{ W}$$

C.O.P. of large cryogenic helium refrigerators



Refrigeration cycles and duties

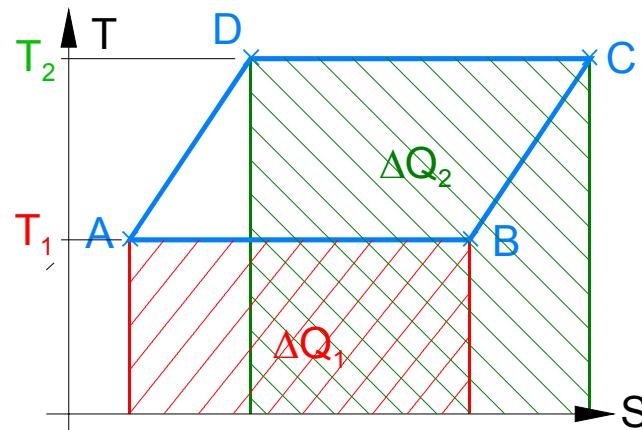
Introduction to the T-S diagram



Thermodynamic transformation from A to B,
if reversible:

$$\Delta Q = \int_A^B T \cdot dS$$

To make a refrigeration cycle, need a substance, the entropy of
which depends on some other variable than temperature



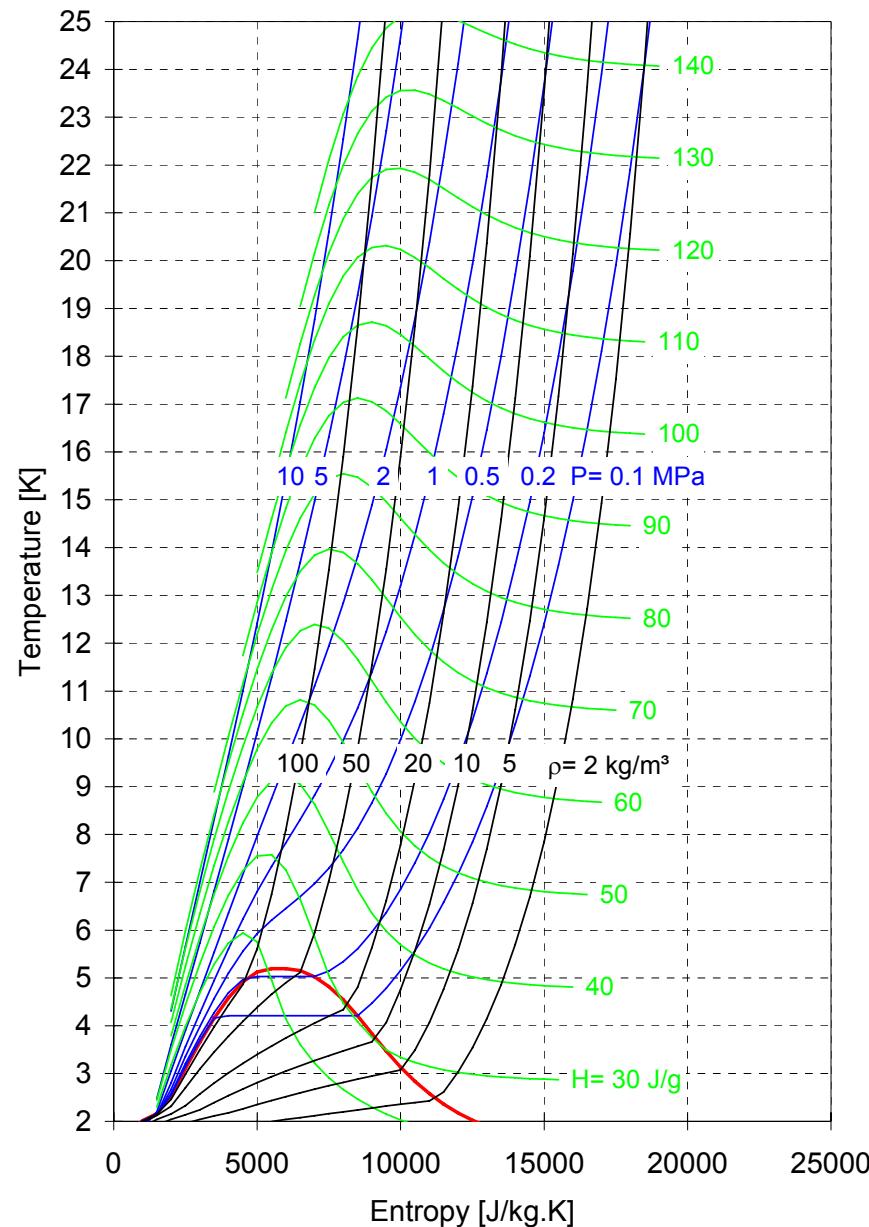
Pressure of gas: Compression/expansion cycle
Magnetization of solid: magnetic refr. cycle

ΔQ_1 : heat absorbed at T_1

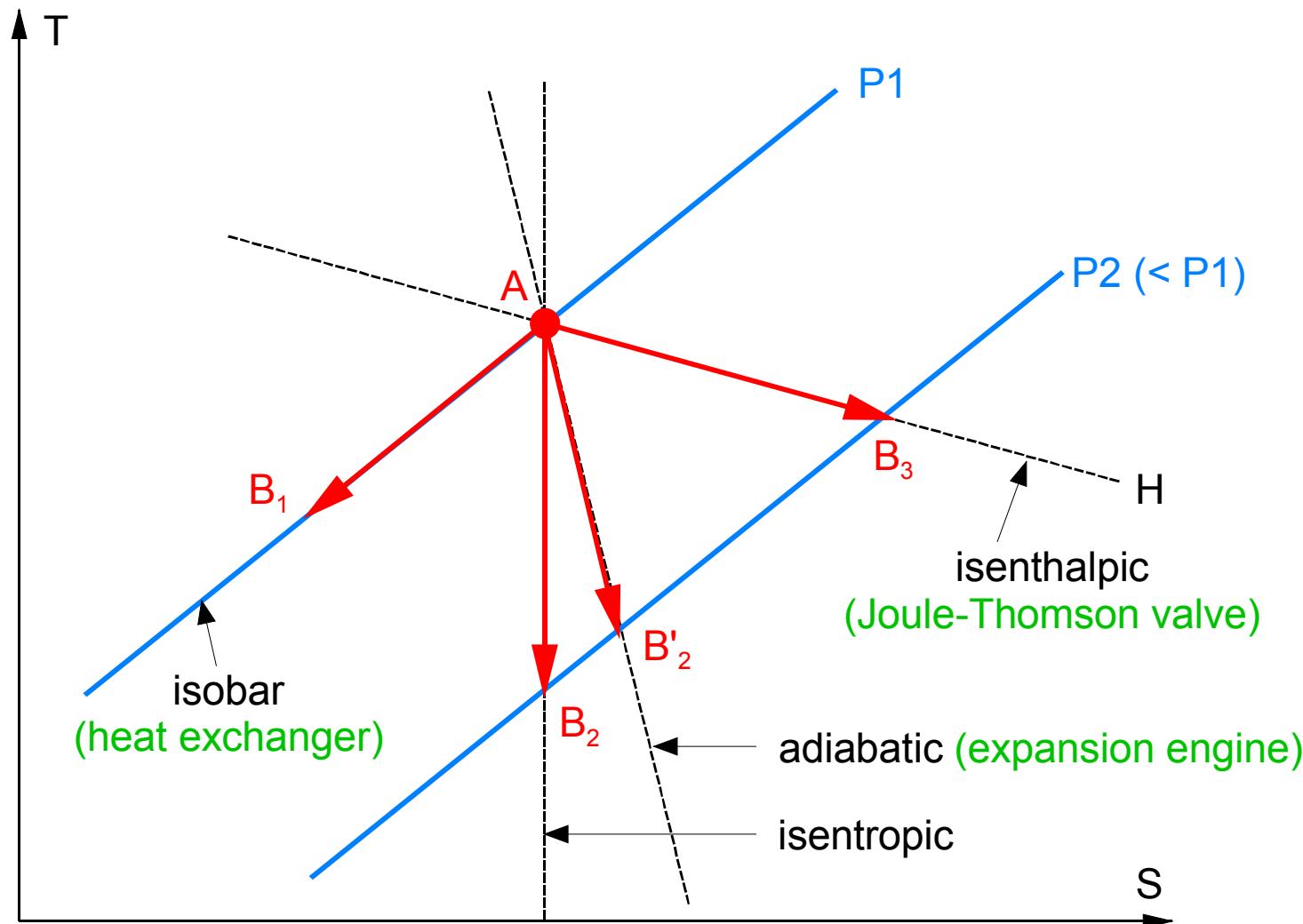
ΔQ_2 : heat rejected at T_2

→ Refrigeration cycle A B C D

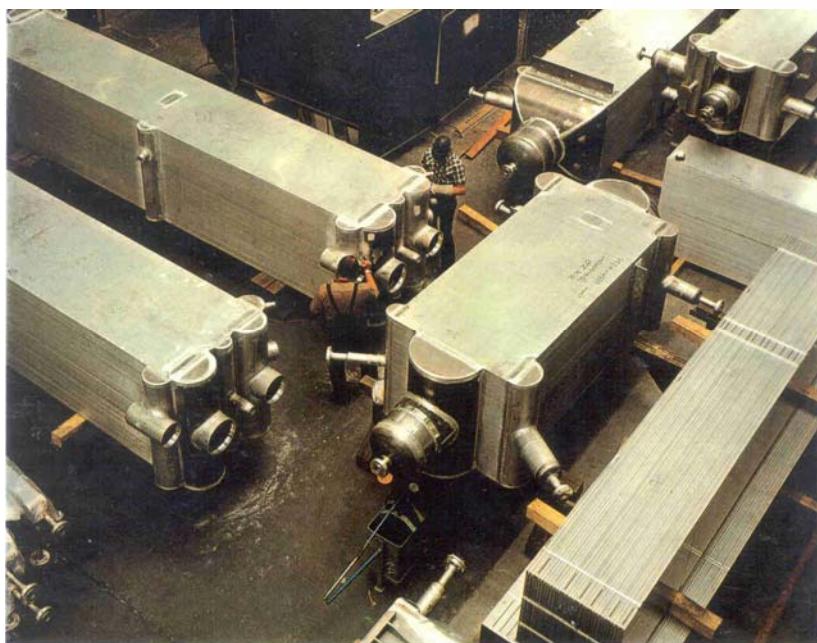
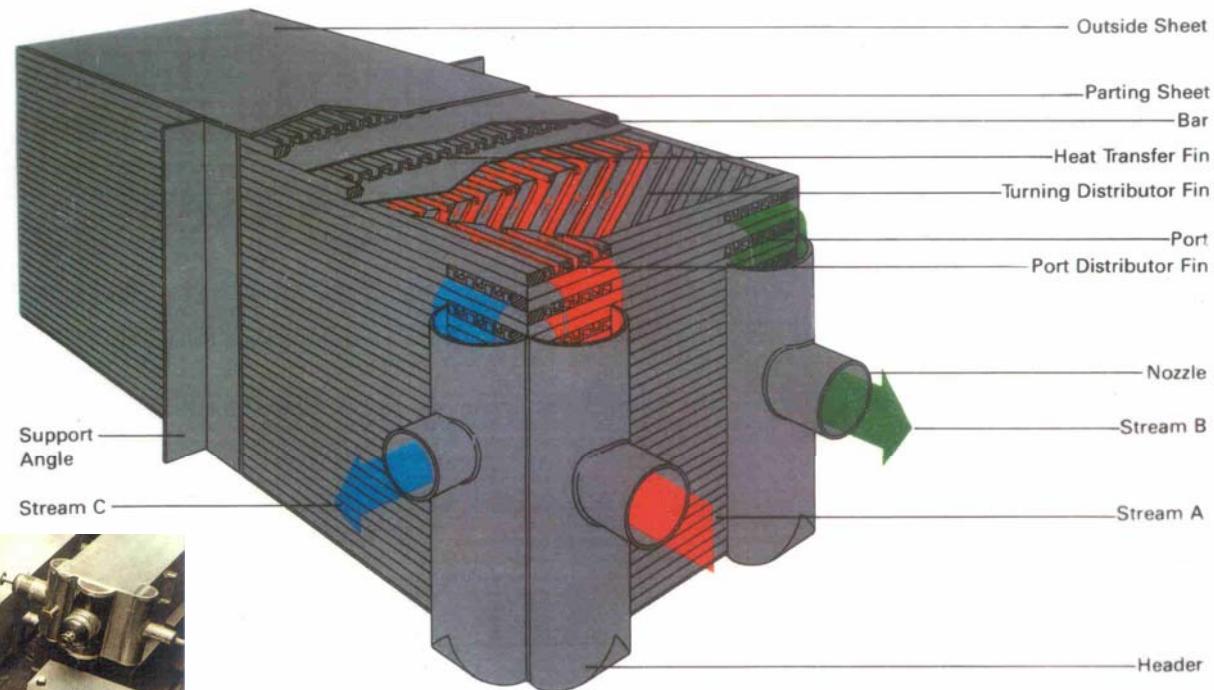
T-S diagram for helium



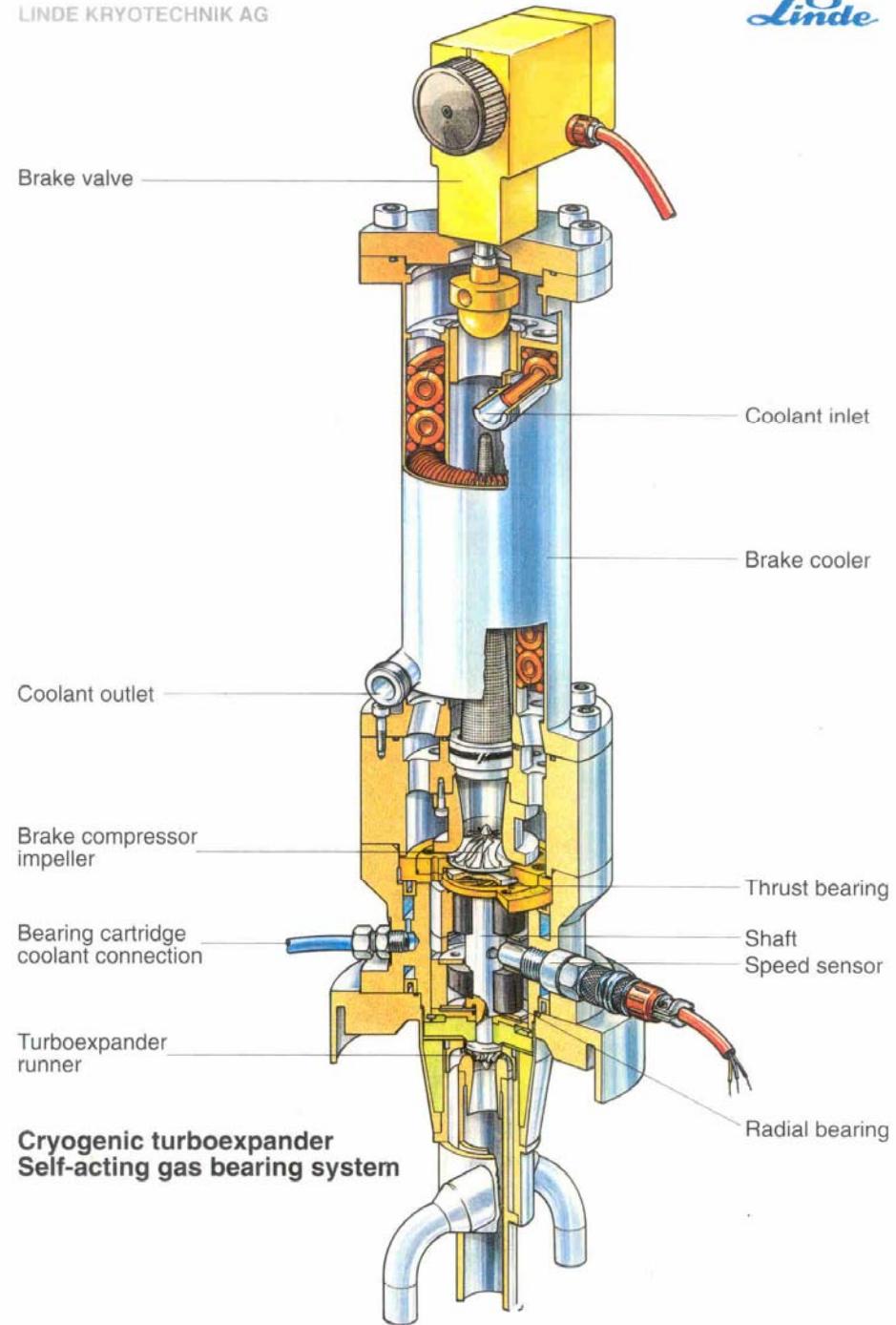
Elementary cooling processes on T-S diagram



Brazed aluminium plate heat exchanger



Cryogenic turbo-expander

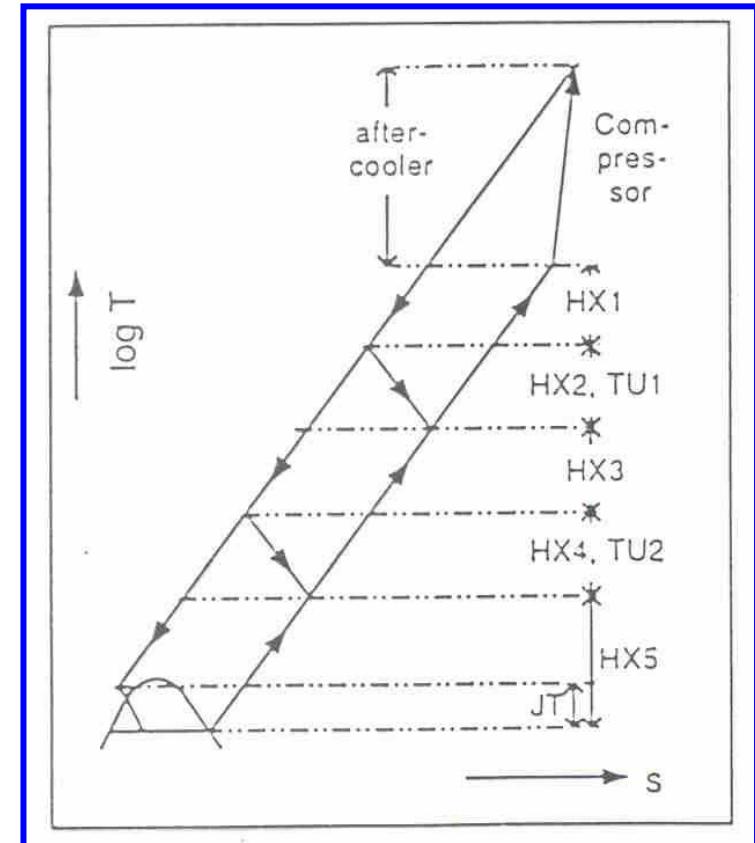
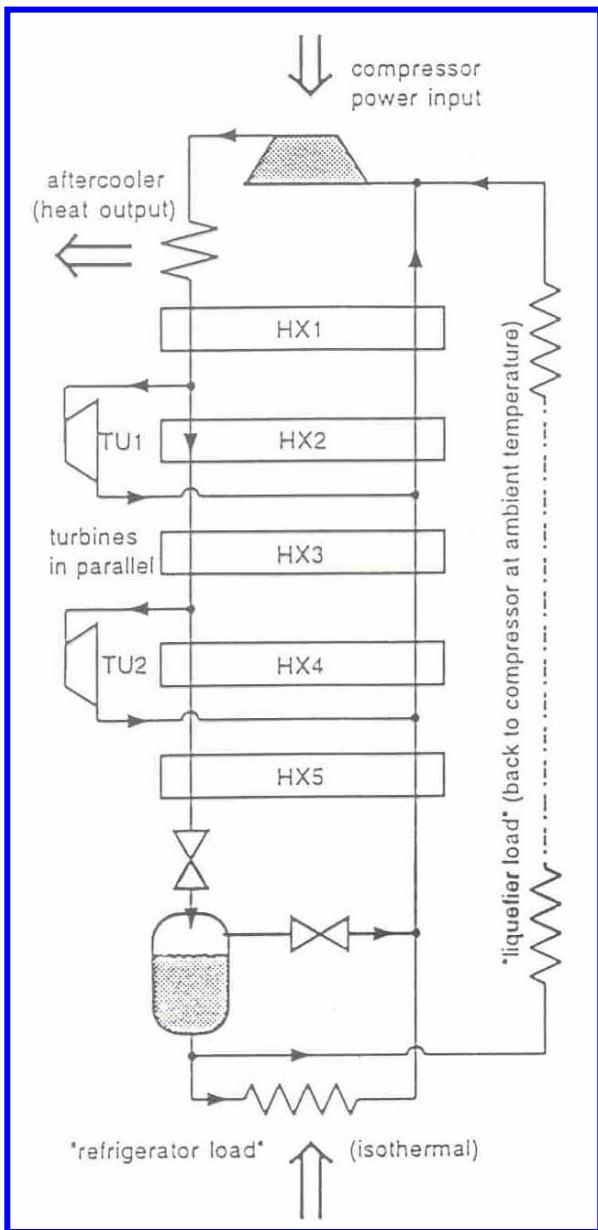


Maximum Joule-Thomson inversion temperatures

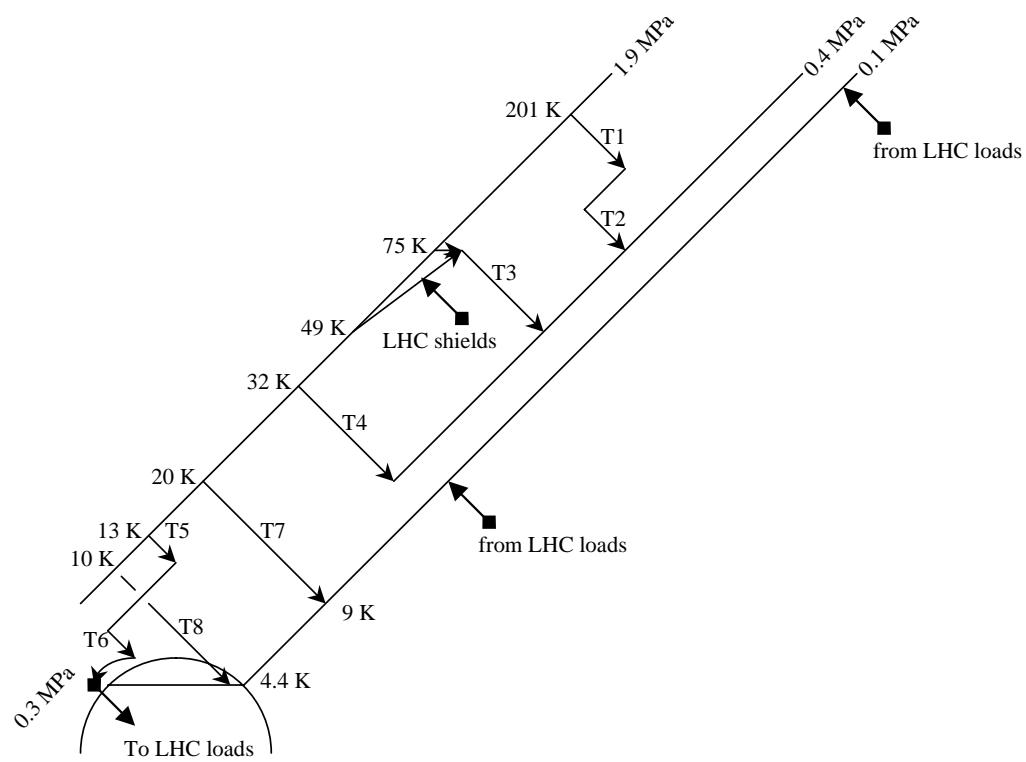
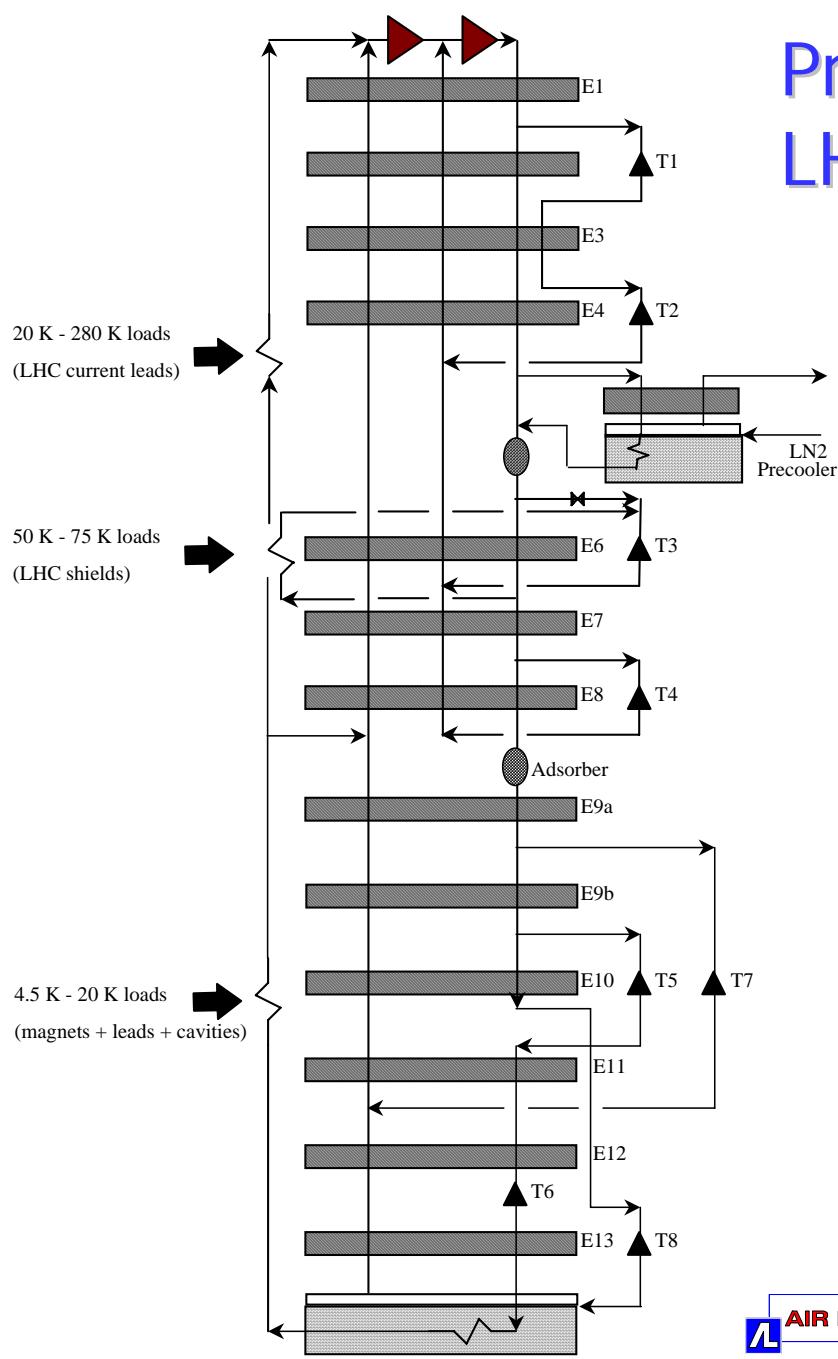
Cryogen	Maximum inversion temperature [K]
Helium	43
Hydrogen	202
Neon	260
Air	603
Nitrogen	623
Oxygen	761

While air can be cooled down and liquefied by JT expansion from room temperature, helium and hydrogen need precooling down to below inversion temperature by heat exchange or work-extracting expansion (e.g. in turbines)

Two-stage Claude cycle



Process cycle & T-S diagram of LHC 18 kW @ 4.5 K cryoplant



LHC 18 kW @ 4.5 K helium cryoplants



33 kW @ 50 K to 75 K
23 kW @ 4.6 K to 20 K
41 g/s liquefaction
4 MW compressor power
C.O.P. 220-230 W/W @ 4.5 K



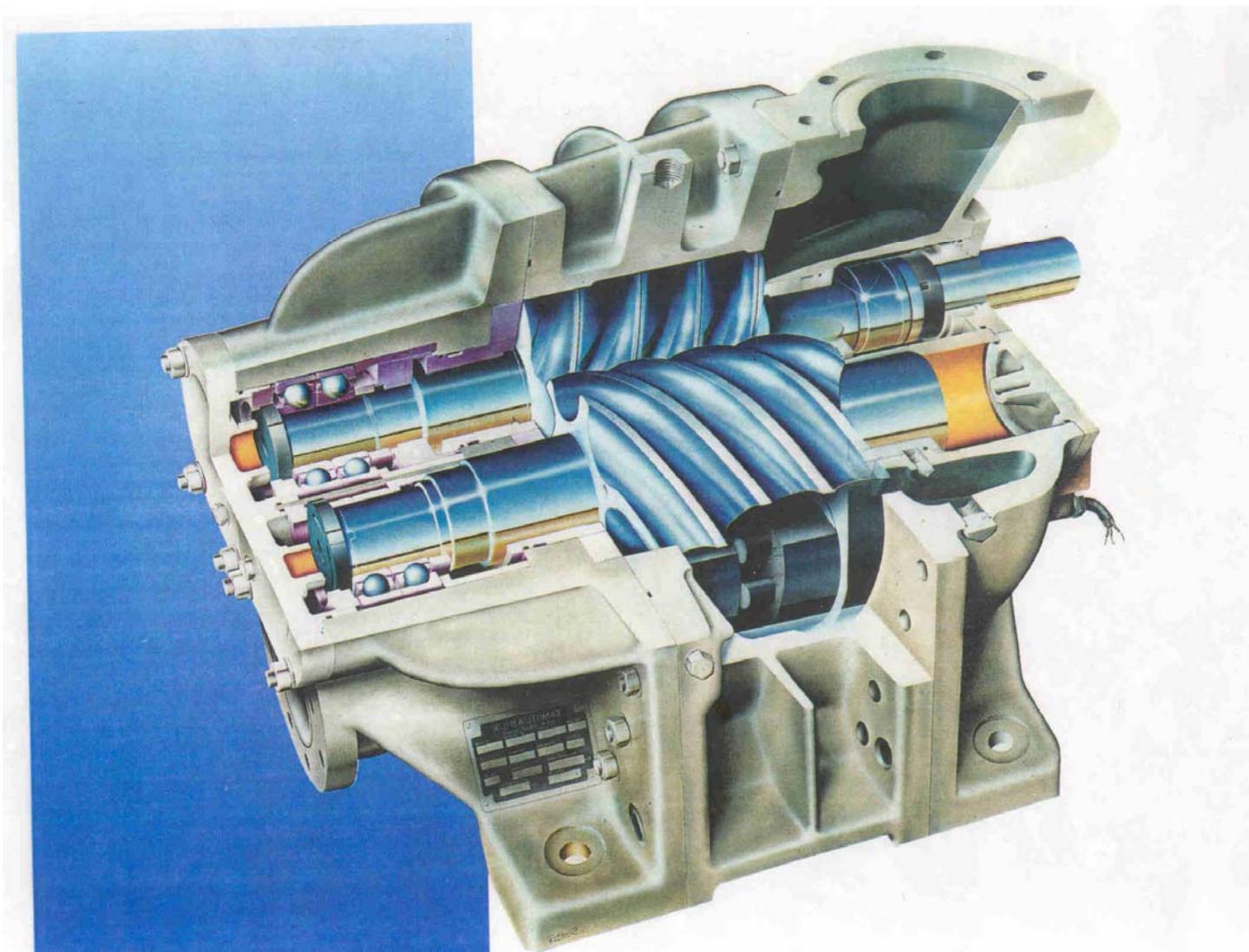
Air Liquide



Linde



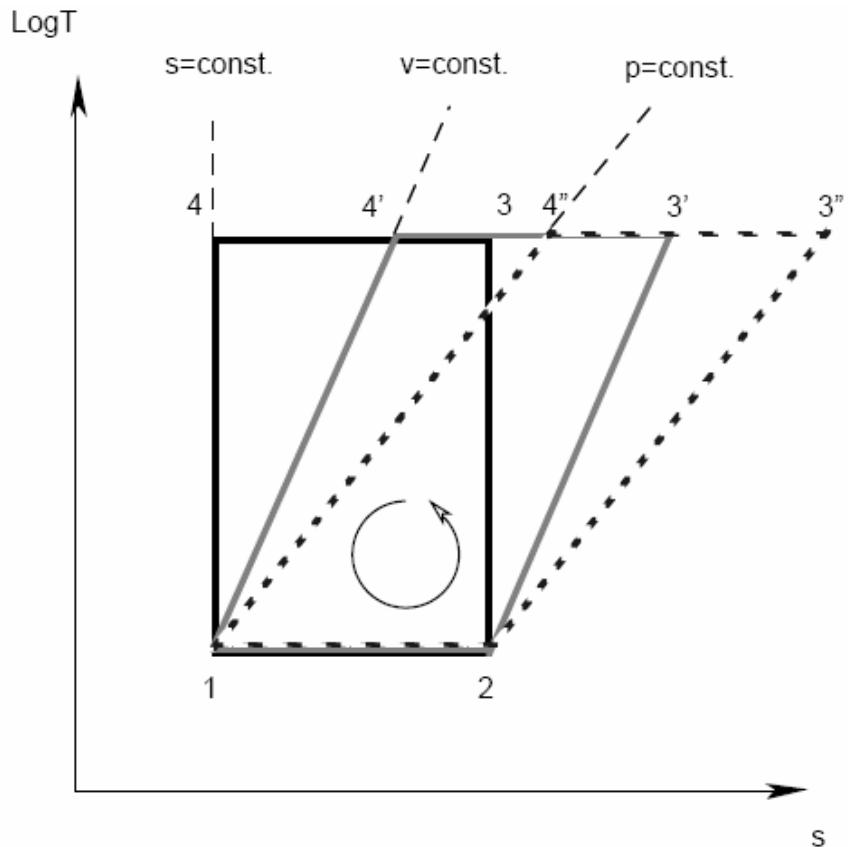
Oil-injected screw compressor



Compressor station of LHC 18 kW@ 4.5 K helium refrigerator

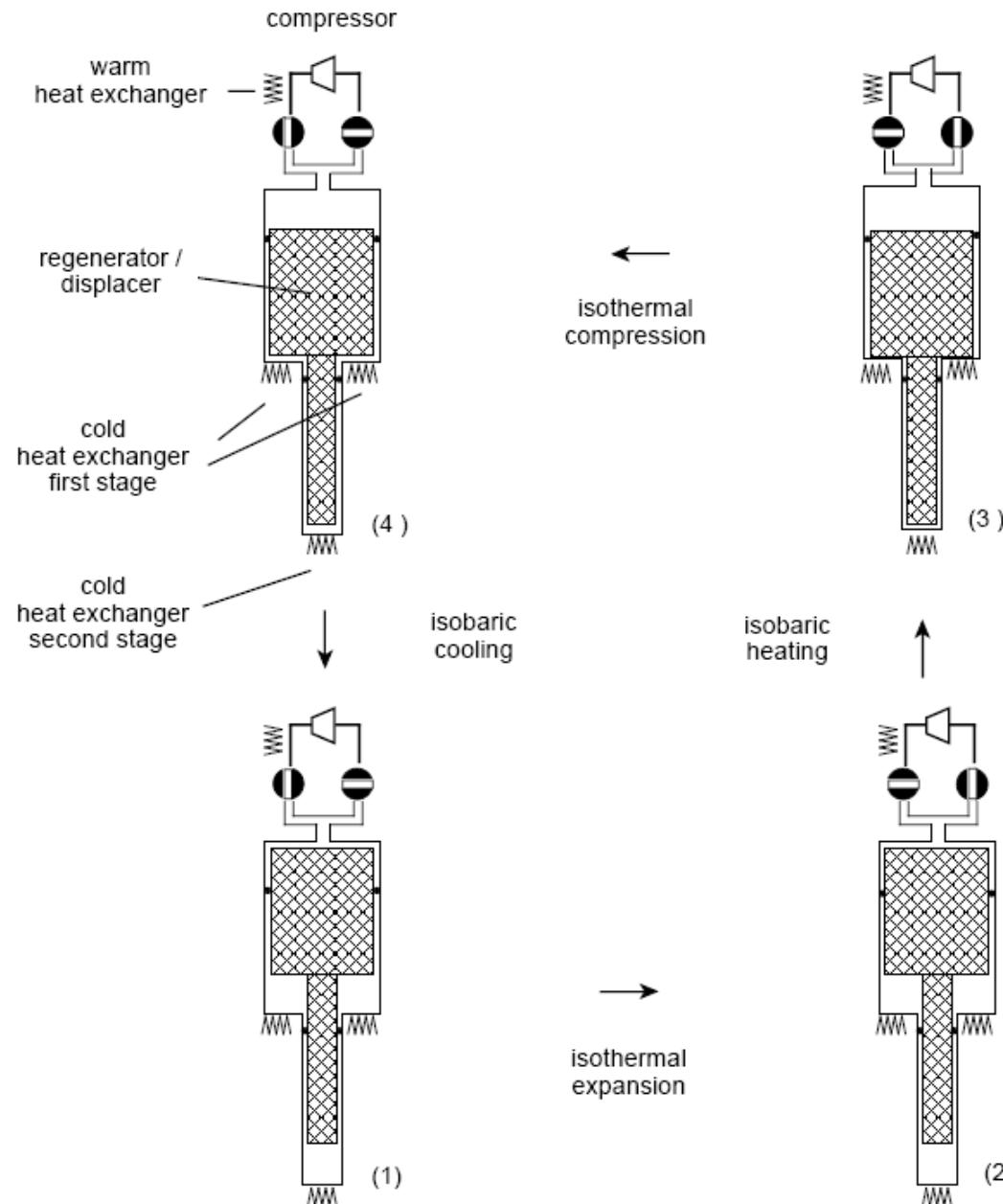


Carnot, Stirling and Ericsson cycles



Carnot cycle (1,2,3,4), Stirling cycle (1,2,3',4') and Ericsson cycle (1,2,3'',4'')

Operation of a Gifford-McMahon cryocooler (Ericsson cycle)



Two-stage Gifford McMahon cryocooler



CRYOMECH PT407 & CP970 compressor
~ 0.7 W @ 4.2 K & 25 W @ 55 K



Contents

- Introduction
- Cryogenic fluids
- Heat transfer & thermal insulation
- Cryogenic distribution & cooling schemes
- Refrigeration & liquefaction
- Cryogen storage & transport
- Thermometry

Specific cost of bulk He storage

Type	Pressure [MPa]	Density [kg/m ³]	Dead volume [%]	Cost [CHF/kg He]
Gas Bag	0.1	0.16	0	300 ⁽¹⁾
MP Vessel	2	3.18	5-25	220-450
HP Vessel	20	29.4	0.5	500 ⁽²⁾
Liquid	0.1	125	13	100-200 ⁽³⁾

(1): Purity non preserved

(2): Not including HP compressors

(3): Not including reliquefier

Bulk helium storage solutions

11000 gallon liquid container



2 MPa gas tanks

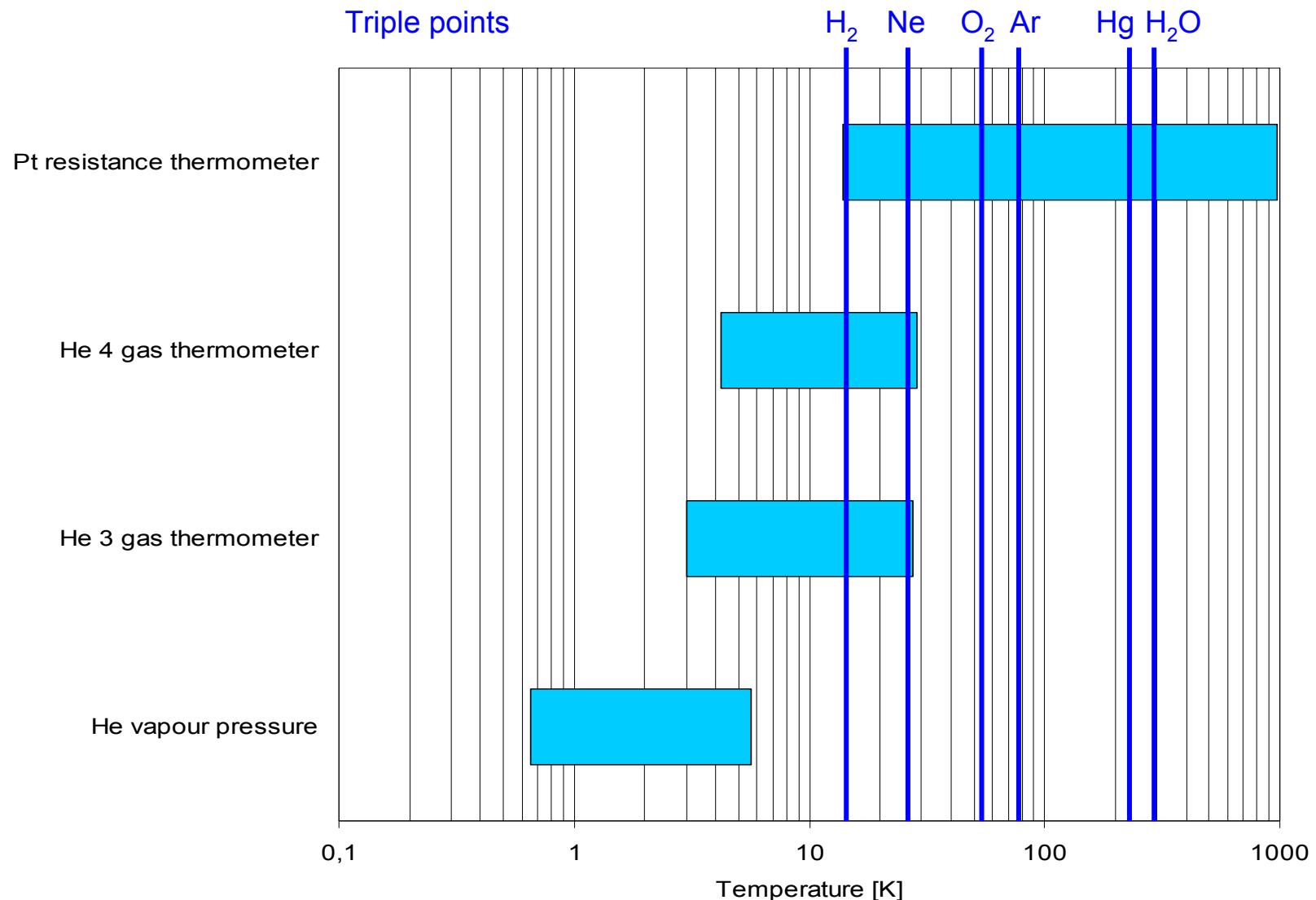


20 MPa gas cylinders

Contents

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Definition of ITS90 in cryogenic range



Primary fixed points of ITS90 in cryogenic range

Fixed point	Temperature [K]
H ₂ triple point	13.8033
Ne triple point	24.5561
O ₂ triple point	54.3584
Ar triple point	83.8058
Hg triple point	234.3156
H ₂ O triple point	273.16 (*)

(*) *exact by definition*

From temperature sensor to practical thermometer



Ge



RhFe wire



RhFe thin film



Cernox



Carbon A-B



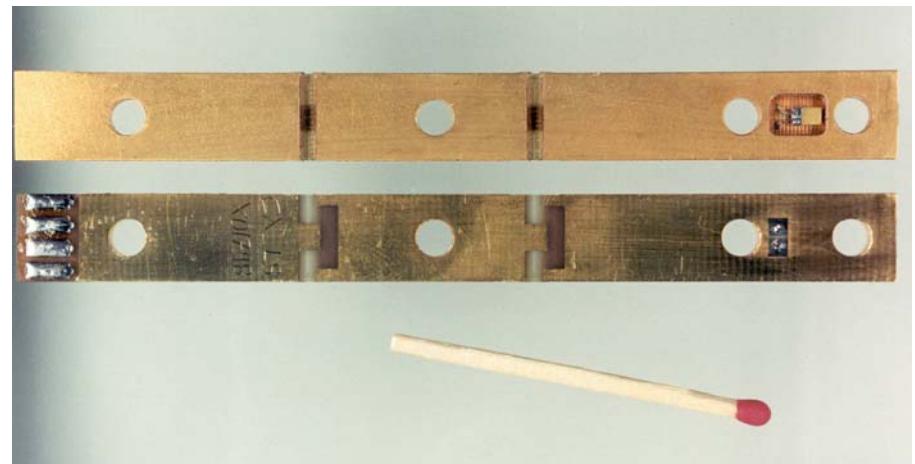
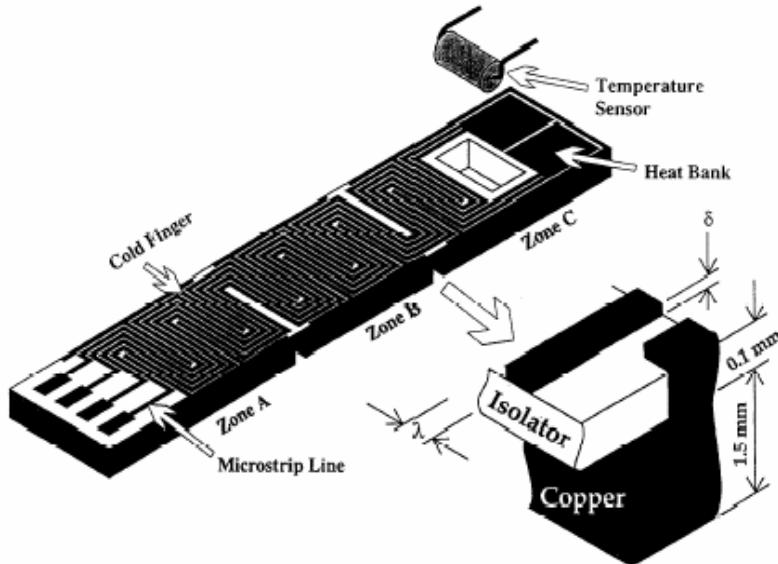
Carbon TVO



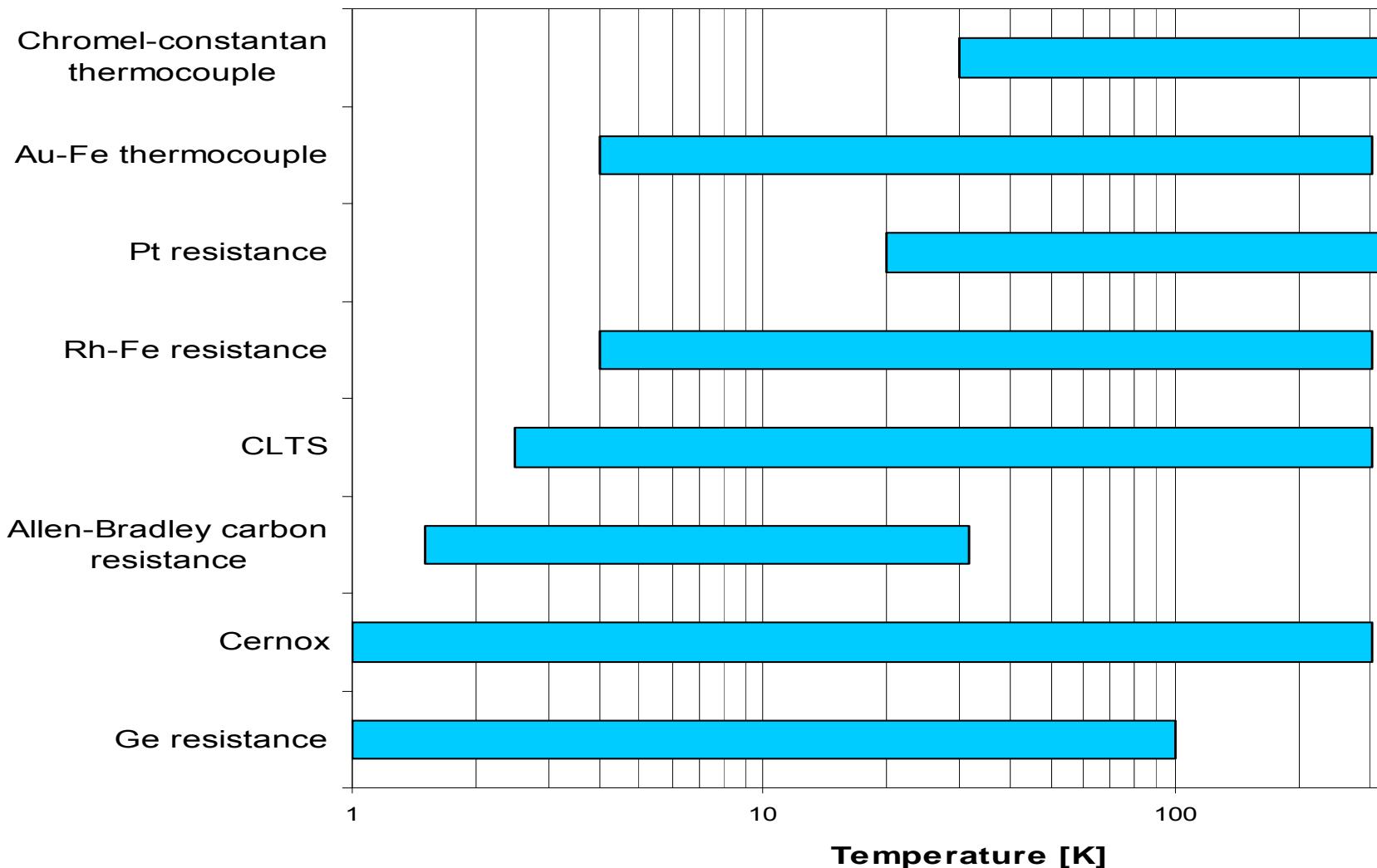
CBT



1cm



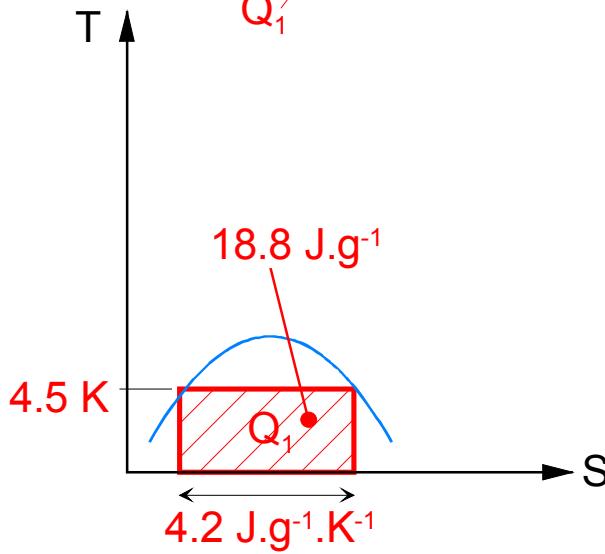
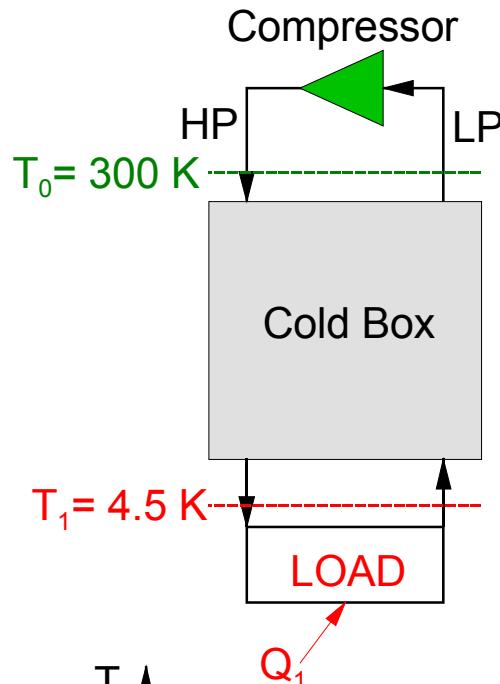
Practical temperature range covered by cryogenic thermometers



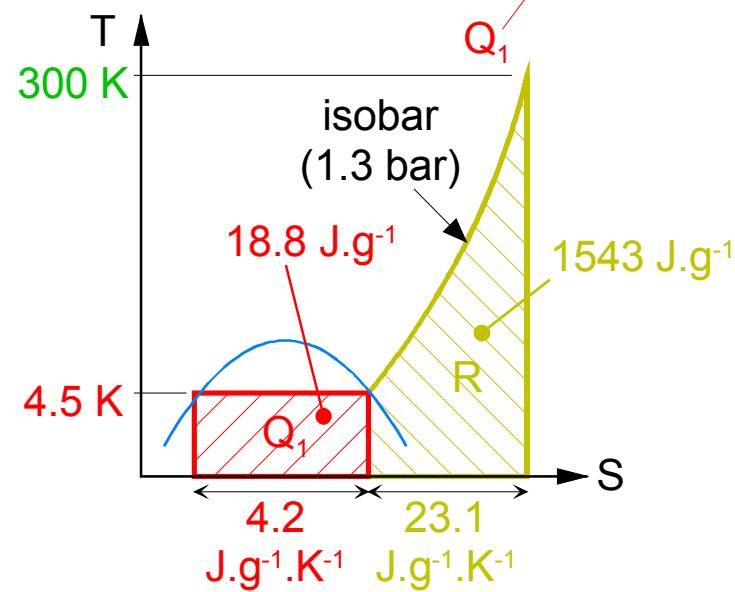
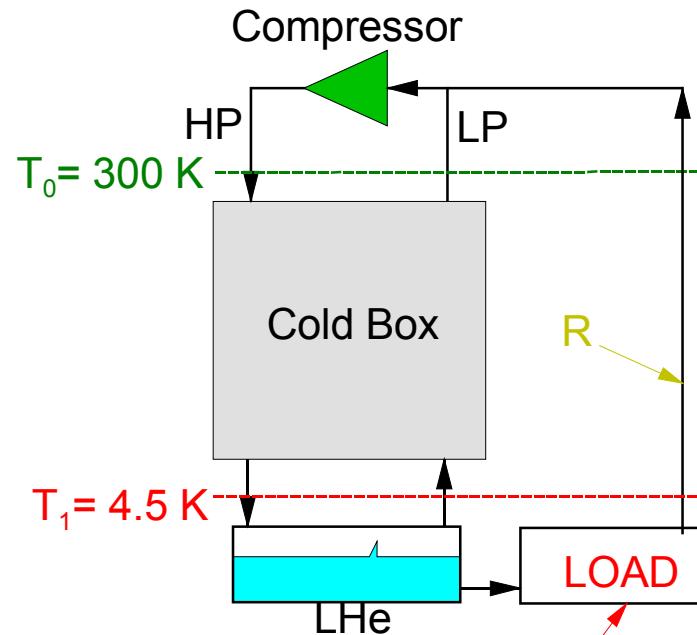
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Refrigerator



Liquefier



Thermodynamic equivalence between refrigeration and liquefaction

What is the isothermal 4.5 K (T_1) refrigeration equivalent to 1 g.s⁻¹ liquefaction of helium?

$$\dot{W}_{\text{min.lique}} = \dot{m}_{\text{lique}} \cdot (T_0 \cdot \Delta S - Q_1 - R)$$

$$\dot{m}_{\text{lique}} = 1 \text{ g.s}^{-1}, T_0 = 300 \text{ K}, \Delta S = 27.3 \text{ J.g}^{-1}.K^{-1}, Q_1 = 18.8 \text{ J.g}^{-1}, R = 1543 \text{ J.g}^{-1}$$

$$\dot{W}_{\text{min.lique}} = 6628 \text{ W}$$

Write that the same amount of work is used to produce isothermal refrigeration at 4.5 K:

$$\dot{W}_{\text{min.refrig}} = \dot{Q}_1 \cdot \left(\frac{T_0}{T_1} - 1 \right)$$

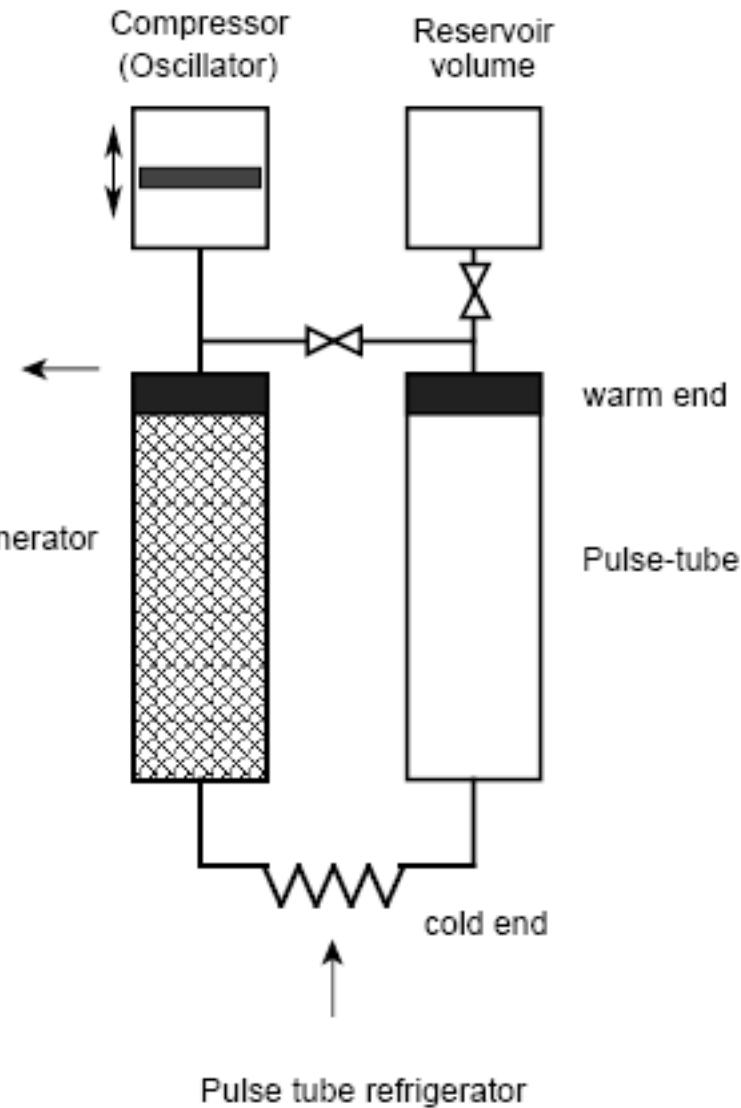
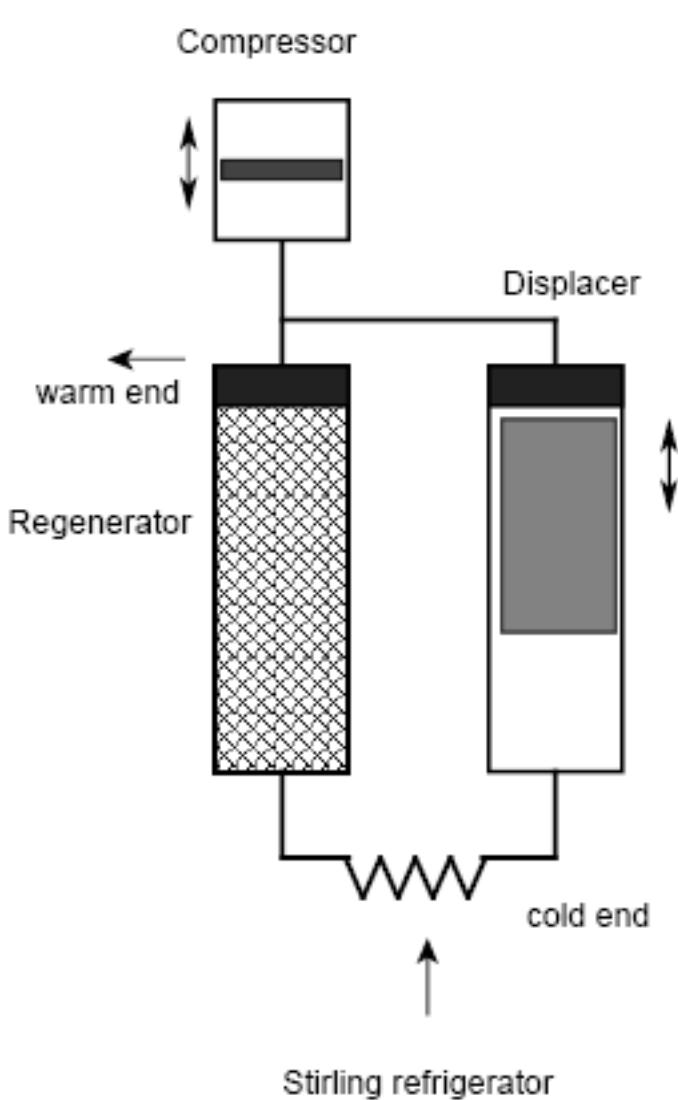
$$\Rightarrow \dot{Q}_1 = 100 \text{ W}$$

$$\dot{W}_{\text{min.refrig}} = \dot{W}_{\text{min.lique}} = 6628 \text{ W}$$

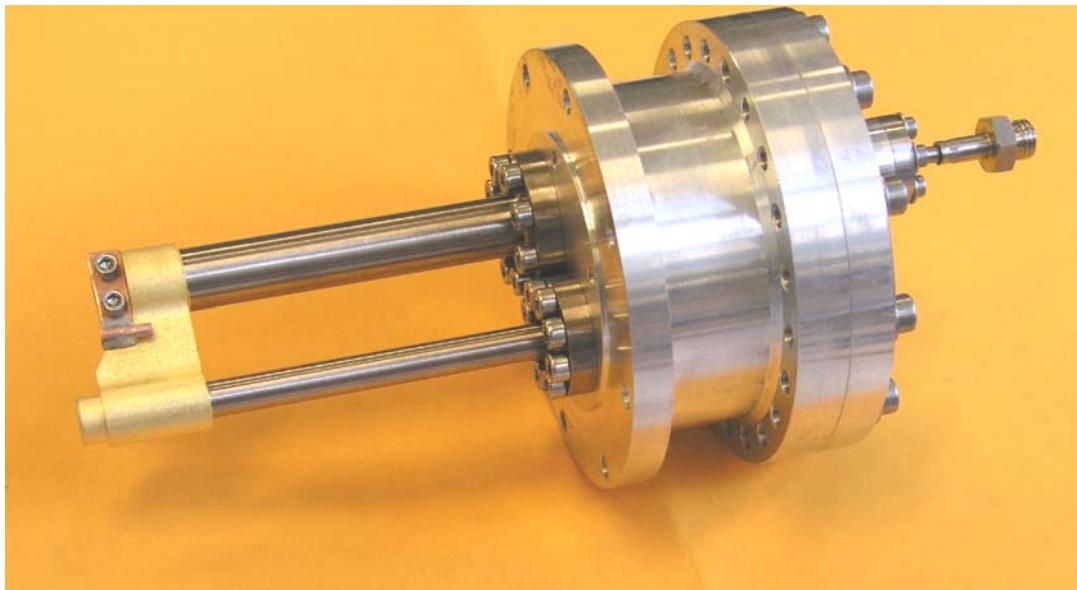
For refrigerators/liquefiers with the same efficiency:

$$1 \text{ g.s}^{-1} \text{ LHe} \equiv 100 \text{ W @ 4.5 K}$$

Stirling and pulse-tube cryocoolers



Mini pulse-tube cryocoolers



ESA MPTC development model – 1W @ 77K



CEA/SBT coaxial PTC – 6W @ 80K