

Formelsammlung

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Contents

I Linear algebra	1
1 Determinant	1
2 linalg:zeug	1
3 Eigenvalues	2
II Geometry	4
4 Trigonometry	4
4.1 Various theorems	4
4.2 Table of values	5
III Calculus	6
4.3 Convolution	6
4.4 Fourier analysis	6
4.4.1 Fourier series	6
4.4.2 Fourier transformation	7
5 List of common integrals	7
IV Probability theory	8
6 Distributions	8
6.0.1 Gauß/Normal distribution	9
6.0.2 Cauchys / Lorentz distribution	9
6.0.3 Binomial distribution	9
6.0.4 Poisson distribution	10
6.0.5 Maxwell-Boltzmann distribution	11
6.1 Central limit theorem	11
V Mechanics	12
7 Lagrange formalism	12
VI Statistical Mechanics	13

8 Entropy	13
VII Thermodynamics	14
9 Processes	14
9.1 Irreversible gas expansion (Gay-Lussac experiment)	14
10 Phase transitions	14
10.0.1 Osmosis	15
10.1 Material properties	15
11 Laws of thermodynamics	16
11.1 Zeroeth law	16
11.2 First law	16
11.3 Second law	17
11.4 Third law	17
12 Ensembles	17
12.1 Potentials	18
13 Ideal gas	18
13.0.1 Molecule gas	19
14 Real gas	20
14.1 Virial expansion	20
14.2 Van der Waals equation	20
15 Ideal quantum gas	21
15.1 Bosons	23
15.2 Fermions	23
15.2.1 Strong degeneracy	24
VIII Electrodynamics	25
16 Maxwell-Equations	25
17 Fields	25
17.1 Electric field	25
17.2 Electric field	25
17.3 Induction	26
18 Hall-Effect	26
18.1 Classical Hall-Effect	26
18.2 Integer quantum hall effect	27
19 Dipole-stuff	28
IX Quantum Mechanics	29
20 Basics	29
20.1 Operators	29
20.2 Probability theory	29
20.2.1 Pauli matrices	30

20.3 Commutator	30
21 Schrödinger equation	31
21.1 Time evolution	31
21.1.1 Schrödinger- and Heisenberg-pictures	32
21.1.2 Ehrenfest theorem	32
21.2 Correspondence principle	32
22 Perturbation theory	33
23 Harmonic oscillator	33
23.1 Creation and Annihilation operators / Ladder operators	34
23.1.1 Harmonischer Oszillator	34
24 Angular momentum	34
24.1 Aharonov-Bohm effect	35
25 Symmetries	35
25.1 Time-reversal symmetry	35
26 Two-level systems (TLS)	35
27 Other	36
28 Hydrogen Atom	36
28.1 Corrections	37
28.1.1 Darwin term	37
28.1.2 Spin-orbit coupling (LS-coupling)	37
28.1.3 Fine-structure	37
28.1.4 Lamb-shift	38
28.1.5 Hyperfine structure	38
28.2 Effects in magnetic field	39
X Condensed matter physics	40
29 Bravais lattice	40
30 Reciprocal lattice	42
30.1 Scattering processes	42
31 Free electron gas	42
31.1 Drude model	43
31.2 Sommerfeld model	43
31.3 2D electron gas	43
31.4 1D electron gas / quantum wire	44
31.5 0D electron gas / quantum dot	44
32 Measurement techniques	44
32.1 ARPES	44
32.2 Scanning probe microscopy SPM	44
33 Fabrication techniques	45
33.1 Epitaxy	45

linalg I

Linear algebra

1 Determinant

2x2 matrix

$$\det \begin{pmatrix} a & b \\ c & d \end{pmatrix} = ad - cb \quad (1)$$

3x3 matrix (Rule of Sarrus)

$$\det \begin{pmatrix} a & b & c \\ d & e & f \\ g & h & i \end{pmatrix} = aei + bfg + cdh - geg - fha - idb \quad (2)$$

Leibniz formula

$$\det(A) = \sum_{\sigma \in S_n} \left(\text{sgn}(\sigma) \prod_{i=1}^n a_{i,\sigma(i)} \right) \quad (3)$$

Product

$$\det(AB) = \det(A)\det(B) \quad (4)$$

Inverse

$$\det(A^{-1}) = \det(A)^{-1} \quad (5)$$

Transposed

$$\det(A^T) = \det(A) \quad (6)$$

2 linalg:zeug

Inverse 2×2 matrix

$$\begin{pmatrix} a & b \\ c & d \end{pmatrix}^{-1} = \frac{1}{ad - bc} \begin{pmatrix} d & -b \\ -c & a \end{pmatrix} \quad (7)$$

Unitary matrix

$$U^\dagger U = \mathbb{1} \quad (8)$$

Singular value decomposition

$$A = U\Lambda V \quad (9)$$

Factorization of complex matrices through rotating
→rescaling →rotation.

A : $m \times n$ matrix, U : $m \times m$ unitary matrix, Λ : $m \times n$ rectangular diagonal matrix with non-negative numbers on the diagonal, V : $n \times n$ unitary matrix

2D rotation matrix

$$R = \begin{pmatrix} \cos \theta & -\sin \theta \\ \sin \theta & \cos \theta \end{pmatrix} \quad (10)$$

3D rotation matrices

$$R_x = \begin{pmatrix} 1 & 0 & 0 \\ 0 & \cos \theta & -\sin \theta \\ 0 & \sin \theta & \cos \theta \end{pmatrix} \quad (11)$$

$$R_y = \begin{pmatrix} \cos \theta & 0 & \sin \theta \\ 0 & 1 & 0 \\ -\sin \theta & 0 & \cos \theta \end{pmatrix} \quad (12)$$

$$R_z = \begin{pmatrix} \cos \theta & -\sin \theta & 0 \\ \sin \theta & \cos \theta & 0 \\ 0 & 0 & 1 \end{pmatrix} \quad (13)$$

Properties of rotation matrices

$$R^T = R^{-1} \quad (14)$$

$$\det R = 1 \quad (15)$$

$$R \in \text{SO}(n) \quad (16)$$

n dimension, $\text{SO}(n)$ special orthogonal group

3 Eigenvalues

Eigenvalue equation

$$Av = \lambda v \quad (17)$$

λ eigenvalue, v eigenvector

Characteristic polynomial
Zeros are the eigenvalues of A

$$\chi_A = \det(A - \lambda \mathbb{1}) \stackrel{!}{=} 0 \quad (18)$$

Kramer's theorem

If H is invariant under T and $|\psi\rangle$ is an eigenstate of H , then $T|\psi\rangle$ is also an eigenstate of H

$$THT^\dagger = H \quad \wedge \quad H|\psi\rangle = E|\psi\rangle \quad \Rightarrow \quad HT|\psi\rangle = ET|\psi\rangle \quad (19)$$

$$A = V\Lambda V^{-1} \quad (20)$$

Eigendecomposition

A diagonalizable, columns of V are eigenvectors v_i , Λ diagonal matrix with eigenvalues λ_i on the diagonal

TODO:Jordan stuff, blockdiagonal matrices, permutations, skalar product lapacescher entwicklungssatz maybe, cramers rule

geo II

Geometry

4 Trigonometry

Exponential function

$$\exp(x) = \sum_{n=0}^{\infty} \frac{x^n}{n!} \quad (21)$$

Sine

$$\sin(x) = \sum_{n=0}^{\infty} (-1)^n \frac{x^{(2n+1)}}{(2n+1)!} \quad (22)$$

$$= \frac{e^{ix} - e^{-ix}}{2i} \quad (23)$$

Cosine

$$\cos(x) = \sum_{n=0}^{\infty} (-1)^n \frac{x^{(2n)}}{(2n)!} \quad (24)$$

$$= \frac{e^{ix} + e^{-ix}}{2} \quad (25)$$

Hyperbolic sine

$$\sinh(x) = -i \sin ix \quad (26)$$

$$= \frac{e^x - e^{-x}}{2} \quad (27)$$

Hyperbolic cosine

$$\cosh(x) = \cos ix \quad (28)$$

$$= \frac{e^x + e^{-x}}{2} \quad (29)$$

4.1 Various theorems

$$1 = \sin^2 x + \cos^2 x \quad (30)$$

Addition theorems

$$\sin(x \pm y) = \sin x \cos y \pm \cos x \sin y \quad (31)$$

$$\cos(x \pm y) = \cos x \cos y \mp \sin x \sin y \quad (32)$$

$$\tan(x \pm y) = \frac{\sin(x \pm y)}{\cos(x \pm y)} = \frac{\tan x \pm \tan y}{1 \mp \tan x \tan y} \quad (33)$$

Double angle

$$\sin 2x = 2 \sin x \cos x \quad (34)$$

$$\cos 2x = \cos^2 x - \sin^2 x = 1 - 2 \sin^2 x \quad (35)$$

$$\tan 2x = \frac{2 \tan x}{1 - \tan^2 x} \quad (36)$$

$$\cos x + b \sin x = \sqrt{1 + b^2} \cos(x - \theta) \quad (37)$$

$$\tan \theta = b$$

4.2 Table of values

Degree	0°	30°	45°	60°	90°	120°	180°	270°
Radian	0	$\frac{\pi}{6}$	$\frac{\pi}{4}$	$\frac{\sqrt{\pi}}{3}$	$\frac{\pi}{2}$	$\frac{2\pi}{3}$	π	$\frac{3\pi}{2}$
$\sin(x)$	0	$\frac{1}{2}$	$\frac{\sqrt{2}}{2}$	$\frac{\sqrt{3}}{2}$	1	$\frac{\sqrt{3}}{2}$	0	-1
$\cos(x)$	1	$\frac{\sqrt{3}}{2}$	$\frac{\sqrt{2}}{2}$	$\frac{1}{2}$	0	$-\frac{1}{2}$	-1	0
$\tan(x)$	0	$\frac{1}{\sqrt{3}}$	$\frac{1}{\sqrt{2}}$	$\frac{1}{2}$	∞	$-\sqrt{3}$	0	∞

cal III

Calculus

4.3 Convolution

Convolution is **commutative**, **associative** and **distributive**.

Definition

$$(f * g)(t) = f(t) * g(t) = \int_{-\infty}^{\infty} f(\tau)g(t - \tau) d\tau \quad (38)$$

Notation

$$f(t) * g(t - t_0) = (f * g)(t - t_0) \quad (39)$$

$$f(t - t_0) * g(t - t_0) = (f * g)(t - 2t_0) \quad (40)$$

Commutativity

$$f * g = g * f \quad (41)$$

Associativity

$$(f * g) * h = f * (g * h) \quad (42)$$

Distributivity

$$f * (g + h) = f * g + f * h \quad (43)$$

Complex conjugate

$$(f * g)^* = f^* * g^* \quad (44)$$

4.4 Fourier analysis

4.4.1 Fourier series

Fourier series
Complex representation

$$f(t) = \sum_{k=-\infty}^{\infty} c_k \exp\left(\frac{2\pi i k t}{T}\right) \quad (45)$$

$$f \in \mathcal{L}^2(\mathbb{R}, \mathbb{C}) \text{ } T\text{-periodic}$$

Fourier coefficients
Complex representation

$$c_k = \frac{1}{T} \int_{-\frac{T}{2}}^{\frac{T}{2}} f(t) \exp\left(-\frac{2\pi i}{T} kt\right) dt \quad \text{for } k \geq 0 \quad (46)$$

$$c_{-k} = \overline{c_k} \quad \text{if } f \text{ real} \quad (47)$$

Fourier series
Sine and cosine representation

$$f(t) = \frac{a_0}{2} + \sum_{k=1}^{\infty} \left(a_k \cos\left(\frac{2\pi}{T} kt\right) + b_k \sin\left(\frac{2\pi}{T} kt\right) \right) \quad (48)$$

$$f \in \mathcal{L}^2(\mathbb{R}, \mathbb{C}) \quad T\text{-periodic}$$

Fourier coefficients
Sine and cosine representation
If f has point symmetry:
 $a_{k>0} = 0$, if f has axial
symmetry: $b_k = 0$

$$a_k = \frac{2}{T} \int_{-\frac{T}{2}}^{\frac{T}{2}} f(t) \cos\left(-\frac{2\pi}{T} kt\right) dt \quad \text{for } k \geq 0 \quad (49)$$

$$b_k = \frac{2}{T} \int_{-\frac{T}{2}}^{\frac{T}{2}} f(t) \sin\left(-\frac{2\pi}{T} kt\right) dt \quad \text{for } k \geq 1 \quad (50)$$

$$a_k = c_k + c_{-k} \quad \text{for } k \geq 0 \quad (51)$$

$$b_k = i(c_k - c_{-k}) \quad \text{for } k \geq 1 \quad (52)$$

TODO:cleanup

4.4.2 Fourier transformation

Fourier transform

$$\hat{f}(k) := \frac{1}{\sqrt{2\pi}^n} \int_{\mathbb{R}^n} e^{-ikx} f(x) dx \quad (53)$$

$$\hat{f} : \mathbb{R}^n \mapsto \mathbb{C}, \quad \forall f \in L^1(\mathbb{R}^n)$$

for $f \in L^1(\mathbb{R}^n)$:

- i) $f \mapsto \hat{f}$ linear in f
- ii) $g(x) = f(x - h) \Rightarrow \hat{g}(k) = e^{-ikh} \hat{f}(k)$
- iii) $g(x) = e^{ih \cdot x} f(x) \Rightarrow \hat{g}(k) = \hat{f}(k - h)$
- iv) $g(\lambda) = f\left(\frac{x}{\lambda}\right) \Rightarrow \hat{g}(k) = \lambda^n \hat{f}(\lambda k)$

5 List of common integrals

Riemann Zeta Function

$$\zeta(s) = \sum_{n=1}^{\infty} \frac{1}{n^s} = \frac{1}{(1 - 2^{(1-s)})\Gamma(s)} \int_0^{\infty} d\eta \frac{\eta^{(s-1)}}{e^{\eta} + 1} \quad (54)$$

pt IV

Probability theory

Mean

$$\langle x \rangle = \int w(x) x \, dx \quad (55)$$

Variance

$$\sigma^2 = (\Delta \hat{x})^2 = \langle \hat{x}^2 \rangle - \langle \hat{x} \rangle^2 = \langle (x - \langle x \rangle)^2 \rangle \quad (56)$$

Standard deviation

$$\sigma = \sqrt{(\Delta x)^2} \quad (57)$$

Median

Value separating lower half
from top half

$$\text{med}(x) = \begin{cases} x_{(n+1)/2} & n \text{ odd} \\ \frac{x_{(n/2)} + x_{((n/2)+1)}}{2} & n \text{ even} \end{cases} \quad (58)$$

x dataset with n elements

Probability density function
Random variable has density
 f . The integral gives the
probability of X taking a
value $x \in [a, b]$.

$$P([a, b]) := \int_a^b f(x) \, dx \quad (59)$$

f normalized: $\int_{-\infty}^{\infty} f(x) \, dx = 1$

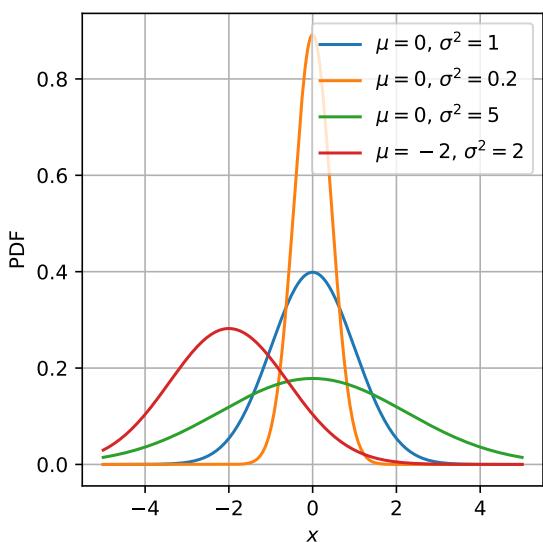
Cumulative distribution
function

$$F(x) = \int_{-\infty}^x f(t) \, dt \quad (60)$$

f probability density function

6 Distributions

6.0.1 Gauß/Normal distribution

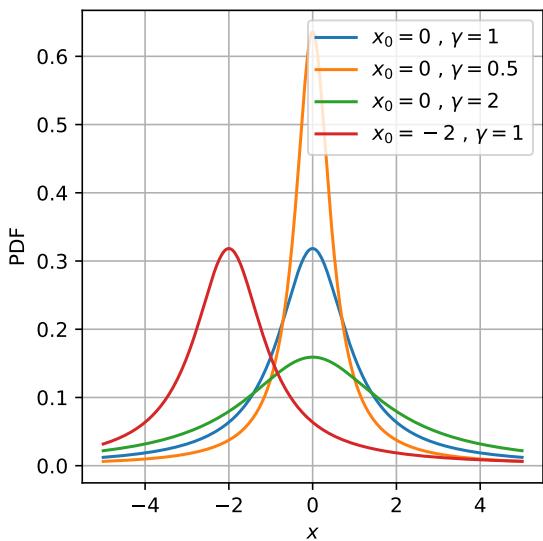


parameters	$\mu \in \mathbb{R}, \quad \sigma^2 \in \mathbb{R}$
support	$x \in \mathbb{R}$
pdf	$\frac{1}{\sqrt{2\pi\sigma^2}} \exp\left(-\frac{(x-\mu)^2}{2\sigma^2}\right)$
cdf	$\frac{1}{2} \left[1 + \operatorname{erf}\left(\frac{x-\mu}{\sqrt{2}\sigma}\right) \right]$
mean	μ
median	μ
variance	σ^2

Density function of the standard normal distribution
 $\mu = 0, \sigma = 1$

$$\varphi(x) = \frac{1}{\sqrt{2\pi}} e^{-\frac{1}{2}x^2} \quad (61)$$

6.0.2 Cauchys / Lorentz distribution

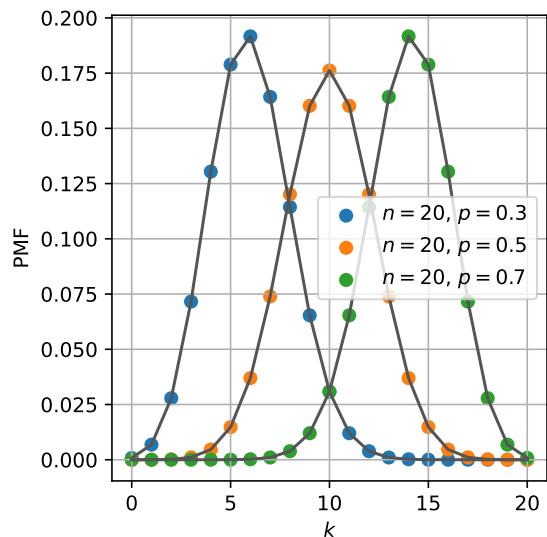


parameters	$x_0 \in \mathbb{R}, \quad \gamma \in \mathbb{R}$
support	$x \in \mathbb{R}$
pdf	$\frac{1}{\pi\gamma \left[1 + \left(\frac{x-x_0}{\gamma} \right)^2 \right]}$
cdf	$\frac{1}{\pi} \arctan\left(\frac{x-x_0}{\gamma}\right) + \frac{1}{2}$
mean	undefined
median	x_0
variance	undefined

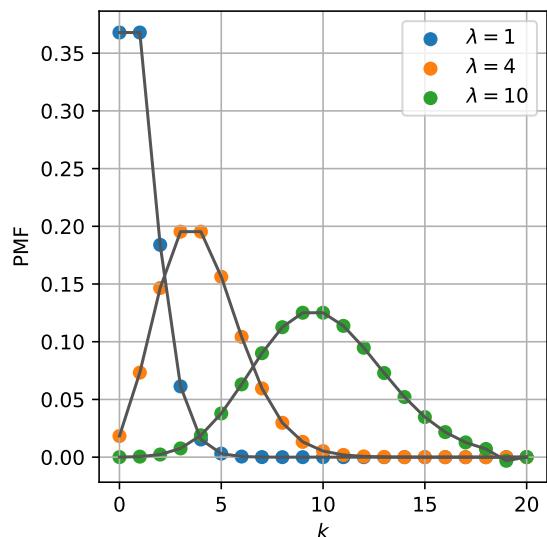
Also known as **Cauchy-Lorentz distribution**, **Lorentz(ian) function**, **Breit-Wigner distribution**.

6.0.3 Binomial distribution

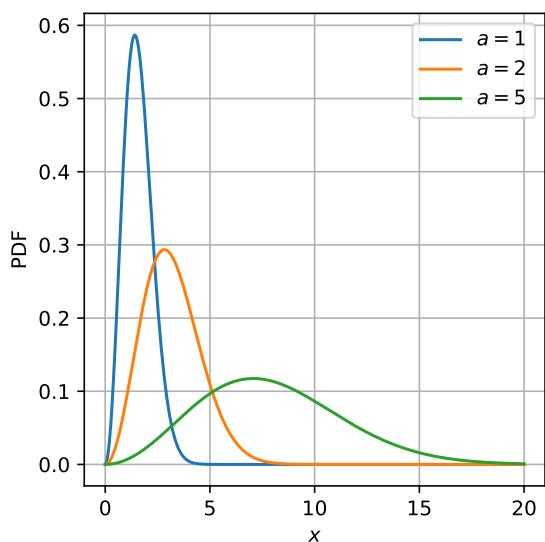
For the number of trials going to infinity ($n \rightarrow \infty$), the binomial distribution converges to the poisson distribution



6.0.4 Poisson distribution



6.0.5 Maxwell-Boltzmann distribution



parameters	$a > 0$
support	$x \in (0, \infty)$
pdf	$\sqrt{\frac{2}{\pi}} \frac{x^2}{a^3} \exp\left(-\frac{x^2}{2a^2}\right)$
cdf	$\text{erf}\left(\frac{x}{\sqrt{2}a}\right) - \sqrt{\frac{2}{\pi}} \frac{x}{a} \exp\left(-\frac{x^2}{2a^2}\right)$
mean	$2a \frac{2}{\pi}$
median	
variance	$\frac{a^2(3\pi - 8)}{\pi}$

6.1 Central limit theorem

Suppose X_1, X_2, \dots is a sequence of independent and identically distributed random variables with $\langle X_i \rangle = \mu$ and $(\Delta X_i)^2 = \sigma^2 < \infty$. As N approaches infinity, the random variables $\sqrt{N}(\bar{X}_N - \mu)$ converge to a normal distribution $\mathcal{N}(0, \sigma^2)$.

That means that the variance scales with $\frac{1}{\sqrt{N}}$ and statements become accurate for large N .

Mechanics

7 Lagrange formalism

The Lagrange formalism is often the most simple approach to get the equations of motion, because with suitable generalized coordinates obtaining the Lagrange function is often relatively easy.

The generalized coordinates are chosen so that the constraints are automatically fulfilled. For example, the generalized coordinate for a 2D pendulum is $q = \varphi$, with $\vec{x} = \begin{pmatrix} \cos \varphi \\ \sin \varphi \end{pmatrix}$.

Lagrange function

$$\mathcal{L} = T - V \quad (62)$$

T kinetic energy, V potential energy

Lagrange equations (2nd type)

$$\frac{d}{dt} \frac{\partial \mathcal{L}}{\partial \dot{q}_i} - \frac{\partial \mathcal{L}}{\partial q_i} = 0 \quad (63)$$

q generalized coordinates

Canonical Momentum

$$p = \frac{\partial \mathcal{L}}{\partial \dot{q}} \quad (64)$$

Hamiltonian

Hamiltonian can be derived from the Lagrangian using a Legendre transformation

$$H(q, p) = p \dot{q} - \mathcal{L}(q, \dot{q}(q, p)) \quad (65)$$

TODO: Legendre trafo

stat VI

Statistical Mechanics

Intensive quantities: Additive for subsystems (system size dependent): $S(\lambda E, \lambda V, \lambda N) = \lambda S(E, V, N)$

Extensive quantities: Independent of system size, ratio of two intensive quantities

Liouville equation

$$\frac{\partial \rho}{\partial t} = - \sum_{i=1}^N \left(\frac{\partial \rho}{\partial q_i} \frac{\partial H}{\partial p_i} - \frac{\partial \rho}{\partial p_i} \frac{\partial H}{\partial q_i} \right) = \{H, \rho\} \quad (66)$$

{ } poisson bracket

8 Entropy

Positive-definite and additive

$$S \geq 0 \quad (67)$$

$$S(E_1, E_2) = S_1 + S_2 \quad (68)$$

Von-Neumann

$$S = -k_B \langle \log \rho \rangle = -k_B \text{tr}(\rho \log \rho) \quad (69)$$

ρ density matrix

Gibbs

$$S = -k_B \sum_n p_n \log p_n \quad (70)$$

p_n probability for micro state n

Boltzmann

$$S = k_B \log \Omega \quad (71)$$

Ω #micro states

Temperature

$$\frac{1}{T} := \left(\frac{\partial S}{\partial E} \right)_V \quad (72)$$

Pressure

$$p = T \left(\frac{\partial S}{\partial V} \right)_E \quad (73)$$

td VII

Thermodynamics

Thermal wavelength

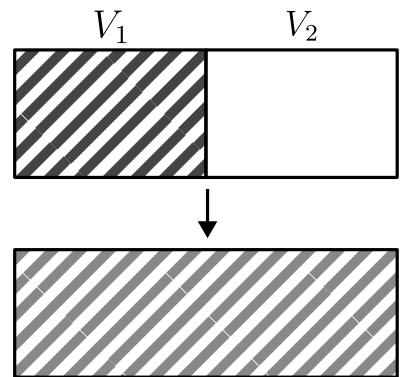
$$\lambda = \frac{h}{\sqrt{2\pi m k_B T}} \quad (74)$$

9 Processes

- **isobaric:** constant pressure $p = \text{const}$
- **isochoric:** constant volume $V = \text{const}$
- **isothermal:** constant temperature $T = \text{const}$
- **isentropic:** constant entropy $S = \text{const}$
- **isenthalpic:** constant enthalpy $H = \text{const}$
- **adiabatic:** no heat transfer $\Delta Q = 0$
- **quasistatic:** happens so slow, the system always stays in td. equilibrium
- **reversible:** reversible processes are always quasistatic and no entropie is created $\Delta S = 0$

9.1 Irreversible gas expansion (Gay-Lussac experiment)

A classical gas in a system with volume V_1 is separated from another system with volume V_2 . In the Gay-Lussac experiment, the separation is removed and the gas flows into V_2 .



Entropy change

$$\Delta S = N k_B \ln \left(\frac{V_1 + V_2}{V_1} \right) > 0 \quad (75)$$

TODO:Reversible

TODO:Quasistatischer T-Ausgleich

TODO:Joule-Thompson Prozess

10 Phase transitions

A phase transition is a discontinuity in the free energy F or Gibbs energy G or in one of their derivatives. The degree of the phase transition is the degree of the derivative which exhibits the discontinuity.

Latent heat

Heat required to bring substance from phase 1 to phase 2

$$Q_L = T\Delta S \quad (76)$$

ΔS entropy change of the phase transition

Clausius-Clapyeron equation
Slope of the coexistence curve

$$\frac{dp}{dT} = \frac{Q_L}{T\Delta V} \quad (77)$$

ΔV Volume change of the phase transition

Phase transition
At the coexistence curve

$$G_1 = G_2 \quad (78)$$

and therefore

$$\mu_1 = \mu_2 \quad (79)$$

Gibbs rule / Phase rule

$$f = c - p + 2 \quad (80)$$

c #components, f #degrees of freedom, p #phases

10.0.1 Osmosis

Osmosis is the spontaneous net movement or diffusion of solvent molecules through a selectively-permeable membrane, which allows through the solvent molecules, but not the solute molecules. The direction of the diffusion is from a region of high water potential (region of lower solute concentration) to a region of low water potential (region of higher solute concentration), in the direction that tends to equalize the solute concentrations on the two sides.

Osmotic pressure

$$p_{\text{osm}} = k_B T \frac{N_c}{V} \quad (81)$$

N_c #dissolved particles

10.1 Material properties

Heat capacity

$$c = \frac{Q}{\Delta T} \quad (82)$$

Q heat

Isochoric heat capacity

$$c_v = \left(\frac{\partial Q}{\partial T} \right)_V = \left(\frac{\partial U}{\partial T} \right)_V \quad (83)$$

U internal energy

Isobaric heat capacity

$$c_p = \left(\frac{\partial Q}{\partial T} \right)_P = \left(\frac{\partial H}{\partial T} \right)_P \quad (84)$$

H enthalpy

Bulk modules

$$K = -V \frac{dp}{dV} \quad (85)$$

p pressure, *V* initial volume

Compressibility

$$\kappa = -\frac{1}{V} \frac{\partial V}{\partial p} \quad (86)$$

Isothermal compressibility

$$\kappa_T = -\frac{1}{V} \left(\frac{\partial V}{\partial p} \right)_T = \frac{1}{K} \quad (87)$$

Adiabatic compressibility

$$\kappa_S = -\frac{1}{V} \left(\frac{\partial V}{\partial p} \right)_S \quad (88)$$

Thermal expansion coefficient

$$\alpha = \frac{1}{V} \left(\frac{\partial V}{\partial T} \right)_{p,N} \quad (89)$$

11 Laws of thermodynamics

11.1 Zeroeth law

If two systems are each in thermal equilibrium with a third, they are also in thermal equilibrium with each other.

$$A \xleftrightarrow{th.eq.} C \quad \wedge \quad B \xleftrightarrow{th.eq.} C \quad \Rightarrow \quad A \xleftrightarrow{th.eq.} B \quad (90)$$

11.2 First law

In a process without transfer of matter, the change in internal energy, ΔU , of a thermodynamic system is equal to the energy gained as heat, Q , less the thermodynamic work, W , done by the system on its surroundings.

Internal energy change

$$\Delta U = \delta Q - \delta W \quad (91)$$

$$dU = T dS - p dV \quad (92)$$

11.3 Second law

Clausius: Heat can never pass from a colder to a warmer body without some other change, connected therewith, occurring at the same time.

Kelvin: It is impossible for a self-acting machine, unaided by any external agency, to convey heat from one body to another at a higher temperature.

11.4 Third law

It is impossible to cool a system to absolute zero.

$$\lim_{T \rightarrow 0} s(T) = 0 \quad (93)$$

and therefore also

Entropy density

$$(94)$$

$$\lim_{T \rightarrow 0} c_V = 0 \quad (95)$$

$$s = \frac{S}{N}$$

12 Ensembles

Table 1: caption

	td:ensembles:mk	td:ensembles:k	td:ensembles:gk
variables	E, V, N	T, V, N	T, V, μ
partition_sum	$\Omega = \sum_n 1$	$Z = \sum_n e^{-\beta E_n}$	$Z_g = \sum_n e^{-\beta(E_n - \mu N_n)}$
probability	$p_n = \frac{1}{\Omega}$	$p_n = \frac{e^{-\beta E_n}}{Z}$	$p_n = \frac{e^{-\beta(E_n - \mu N_n)}}{Z_g}$
td_pot	$S = k_B \ln \Omega$	$F = -k_B T \ln Z$	$\Phi = -k_B T \ln Z$
pressure	$p = T \left(\frac{\partial S}{\partial V} \right)_{E,N}$	$p = - \left(\frac{\partial F}{\partial V} \right)_{T,N}$	$p = - \left(\frac{\partial \Phi}{\partial V} \right)_{T,\mu} = -\frac{\Phi}{V}$
entropy	$S = k_B = \ln \Omega$	$S = - \left(\frac{\partial F}{\partial T} \right)_{V,N}$	$S = - \left(\frac{\partial \Phi}{\partial T} \right)_{V,\mu}$

Ergodic hypothesis

Over a long period of time, all accessible microstates in the phase space are equiprobable

$$\langle A \rangle_{\text{Time}} = \langle A \rangle_{\text{Ensemble}} \quad (96)$$

A Observable

12.1 Potentials

Internal energy

$$dU(S, V, N) = T dS - p dV + \mu dN \quad (97)$$

Enthalpy

$$dH(S, p, N) = T dS + V dp + \mu dN \quad (98)$$

Gibbs energy

$$dG(T, p, N) = -S dT + V dp + \mu dN \quad (99)$$

Free energy / Helmholtz energy

$$dF(T, V, N) = -S dT - p dV + \mu dN \quad (100)$$

Grand canonical potential

$$d\Phi(T, V, \mu) = -S dT - p dV - N d\mu \quad (101)$$

TODO:Maxwell Relationen, TD Quadrat

13 Ideal gas

The ideal gas consists of non-interacting, undifferentiable particles.

Phase space volume

$3N$ sphere

$$\Omega(E) = \int_V d^3q_1 \dots \int_V d^3q_N \int d^3p_1 \dots \int d^3p_N \frac{1}{N! h^{3N}} \Theta\left(E - \sum_i \frac{\vec{p}_i^2}{2m}\right) \quad (102)$$

$$= \left(\frac{V}{N}\right)^N \left(\frac{4\pi m E}{3h^2 N}\right)^{\frac{3N}{2}} e^{\frac{5N}{2}} \quad (103)$$

N #particles, h^{3N} volume of a microstate, $N!$ particles are undifferentiable

Entropy

$$S = \frac{5}{2} N k_B + N k_B \ln\left(\frac{V}{N} \left(\frac{2\pi m E}{3h^2 N}\right)^{\frac{3}{2}}\right) \quad (104)$$

Ideal gas equation

$$pV = nRT \quad (105)$$

$$= Nk_B T \quad (106)$$

Equation of state

$$U = \frac{3}{2} N k_B T \quad (107)$$

Equipartition theorem

Each degree of freedom contributes U_D (for classical particle systems)

$$U_D = \frac{1}{2} k_B T \quad (108)$$

Maxwell velocity distribution

See also 6.0.5

$$w(v) dv = 4\pi \left(\frac{\beta m}{2\pi} \right)^{\frac{3}{2}} v^2 e^{-\frac{\beta mv^2}{2}} dv \quad (109)$$

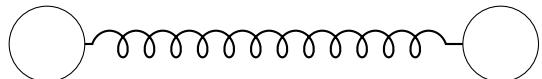
Average quadratic velocity per particle in a 3D gas

$$\langle v^2 \rangle = \int_0^\infty dv v^2 w(v) = \frac{3k_B T}{m} \quad (110)$$

13.0.1 Molecule gas

Molecule gas

2 particles of mass M connected by a “spring” with distance L



Translation

$$p_i = \frac{2\pi\hbar}{L} n_i \quad (111)$$

$$E_{\text{kin}} = \frac{\vec{p}_r^2}{2M} \quad (112)$$

$$n_i \in \mathbb{N}_0, i = x, y, z$$

Vibration

$$E_{\text{vib}} = \hbar\omega \left(n + \frac{1}{2} \right) \quad (113)$$

$$n \in \mathbb{N}_0$$

Rotation

$$E_{\text{rot}} = \frac{\hbar^2}{2I} j(j+1) \quad (114)$$

$$j \in \mathbb{N}_0$$

TODO:Diagram für verschiedene Temperaturen, Weiler Skript p.83

14 Real gas

14.1 Virial expansion

Expansion of the pressure p in a power series of the density ρ .

Virial expansion

The 2nd and 3^d virial coefficient are tabulated for many substances

$$p = k_B T \rho [1 + B(T) \rho + C(T) \rho^2 + \dots] \quad (115)$$

B and C 2nd and 3^d virial coefficient, $\rho = \frac{N}{V}$

Mayer function

$$f(\vec{r}) = e^{-\beta V(i,j)} - 1 \quad (116)$$

$V(i,j)$ pair potential

Second virial coefficient

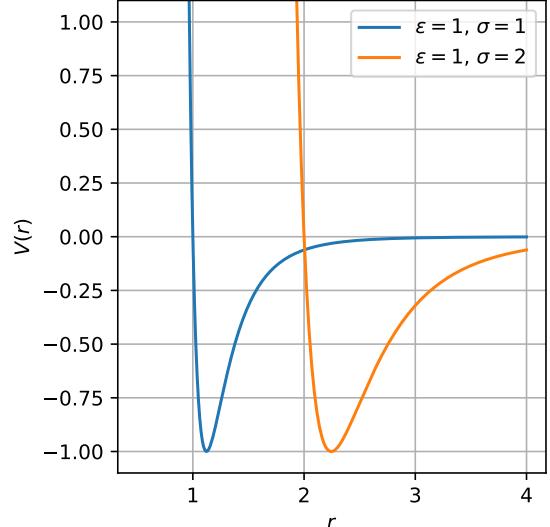
Depends on pair potential between two molecules

$$B = -\frac{1}{2} \int_V d^3 \vec{r} f(\vec{r}) \quad (117)$$

Lennard-Jones potential

Potential between two molecules. Attractive for $r > \sigma$, repulsive for $r < \sigma$

$$V(r) = 4\epsilon \left[\left(\frac{\sigma}{r} \right)^{12} - \left(\frac{\sigma}{r} \right)^6 \right] \quad (118)$$



14.2 Van der Waals equation

Assumes a hard-core potential with a weak attraction.

Partition sum

$$Z_N = \frac{(V - V_0)^N}{\lambda^{3N} N!} e^{\frac{\beta N^2 a}{V}} \quad (119)$$

a internal pressure

Van der Waals equation

$$p = \frac{Nk_B T}{V - b} - \frac{N^2 a}{V^2} \quad (120)$$

b co-volume?

TODO:sometimes N is included in a, b

15 Ideal quantum gas

Fugacity

$$z = e^{\mu\beta} = e^{\frac{\mu}{k_B T}} \quad (121)$$

Occupation number

$$\sum_r n_r = N \quad (122)$$

r states

Undifferentiable particles

$$|p_1, p_2, \dots, p_N\rangle = |p_1\rangle |p_2\rangle \dots |p_N\rangle \quad (123)$$

p_i state

Applying the parity operator yields a *symmetric* (Bosons) and a *antisymmetric* (Fermions) solution

$$\hat{P}_{12}\psi(p_i(\vec{r}_1), p_j(\vec{r}_2)) = \pm\psi(p_i(\vec{r}_1), p_j(\vec{r}_2)) \quad (124)$$

\hat{P}_{12} parity operator swaps 1 and 2, $\pm:$ _{fer}^{bos}

Spin degeneracy factor

$$g_s = 2s + 1 \quad (125)$$

s spin

Density of states

$$g(\epsilon) = g_s \frac{V}{4\pi} \left(\frac{2m}{\hbar^2} \right)^{\frac{3}{2}} \sqrt{\epsilon} \quad (126)$$

g_s td:id_qgas:spin_degeneracy_factor

Occupation number per energy

$$n(\epsilon) d\epsilon = \frac{g(\epsilon)}{e^{\beta(\epsilon-\mu)} \mp 1} d\epsilon \quad (127)$$

td:id_qgas:dos, $\pm:$ _{fer}^{bos}

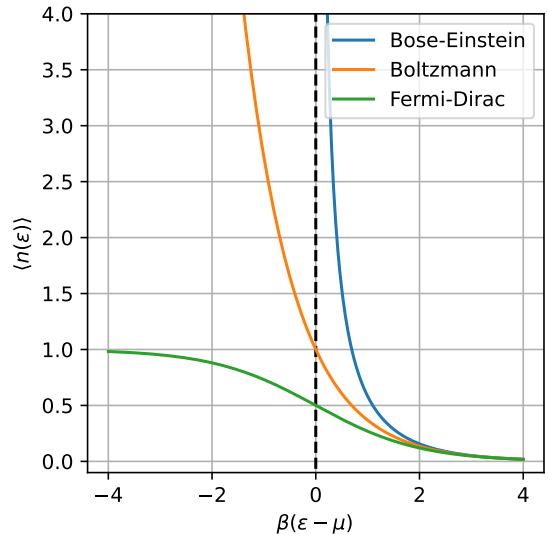
Occupation number

$$\langle n(\epsilon) \rangle = \frac{1}{e^{\beta(\epsilon-\mu)} \mp 1} \quad (128)$$

for $\epsilon - \mu \gg k_B T$

$$= \frac{1}{e^{\beta(\epsilon-\mu)}} \quad (129)$$

$\pm:$ _{bos}
fer



Number of particles

$$\langle N \rangle = \int_0^\infty n(\epsilon) d\epsilon \quad (130)$$

Energy

Equal to the classical ideal gas

$$\langle E \rangle = \int_0^\infty \epsilon n(\epsilon) d\epsilon = \frac{3}{2} p V \quad (131)$$

Equation of state

Bosons: decreased pressure, they like to cluster

Fermions: increased pressure because of the Pauli principle

$$pV = k_B T \ln Z_g \quad (132)$$

after Virial expansion

$$= N k_B T \left[1 \mp \frac{\lambda^3}{2^{5/2} g v} + \mathcal{O}\left(\left(\frac{\lambda^3}{v}\right)^2\right) \right] \quad (133)$$

$\pm:$ _{bos}
fer , $v = \frac{V}{N}$ specific volume

Relevance of qm. corrections

Corrections become relevant when the particle distance is in the order of the thermal wavelength

$$\left(\frac{V}{N}\right)^{\frac{1}{3}} \sim \frac{\lambda}{g_s^{\frac{1}{3}}} \quad (134)$$

Generalized zeta function

$$\left. \begin{aligned} g_\nu(z) \\ f_\nu(z) \end{aligned} \right\} := \frac{1}{\Gamma(\nu)} \int_0^\infty dx \frac{x^{\nu-1}}{e^x z^{-1} \mp 1} \quad (135)$$

15.1 Bosons

Partition sum

$$Z_g = \prod_p \frac{1}{1 - e^{-\beta(\epsilon_p - \mu)}} \quad (136)$$

$$p \in \mathbb{N}_0$$

Occupation number
Bose-Einstein distribution

$$\langle n_p \rangle = \frac{1}{e^{\beta(\epsilon - \mu)} - 1} \quad (137)$$

15.2 Fermions

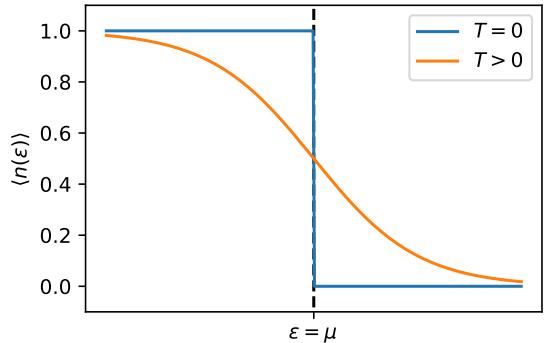
Partition sum

$$Z_g = \prod_p \left(1 + e^{-\beta(\epsilon_p - \mu)} \right) \quad (138)$$

$$p = 0, 1$$

Occupation number
Fermi-Dirac distribution. At $T = 0$ *Fermi edge* at $\epsilon = \mu$

$$\langle n_p \rangle = \frac{1}{e^{\beta(\epsilon - \mu)} + 1} \quad (139)$$



Slater determinant

$$\psi(\vec{r}_1, \vec{r}_2, \dots, \vec{r}_N) = \frac{1}{\sqrt{N!}} \begin{vmatrix} p_1(\vec{r}_1) & p_2(\vec{r}_1) & \dots & p_N(\vec{r}_1) \\ p_1(\vec{r}_2) & p_2(\vec{r}_2) & \dots & p_N(\vec{r}_2) \\ \vdots & \vdots & \ddots & \vdots \\ p_1(\vec{r}_N) & p_2(\vec{r}_N) & \dots & p_N(\vec{r}_N) \end{vmatrix} \quad (140)$$

Fermi energy

$$\epsilon_F := \mu(T = 0) \quad (141)$$

Fermi temperature

$$T_F := \frac{\epsilon_F}{k_B} \quad (142)$$

Fermi impulse

Radius of the *Fermi sphere* in impulse space. States with p_F are in the *Fermi surface*

$$p_F = \hbar k_F = (2mE_F)^{\frac{1}{2}} \quad (143)$$

Specific density

$$v = \frac{N}{V} = \frac{g}{\lambda^3} f_{3/2}(z) \quad (144)$$

f td:id_qgas:generalized_zeta , *g* degeneracy factor, *z* td:id_qgas:fugacity

15.2.1 Strong degeneracy

Sommerfeld expansion
for low temperatures $T \ll T_F$

$$f_\nu(z) = \frac{(\ln z)^\nu}{\Gamma(\nu + 1)} \left(1 + \frac{\pi^6}{6} \frac{\nu(\nu - 1)}{(\ln z)^2} + \dots \right) \quad (145)$$

Energy density

$$\frac{E}{V} = \frac{3}{2} \frac{g}{\lambda^3} k_B T f_{5/2}(z) \quad (146)$$

td:id_qgas:fer:degenerate:sommerfeld

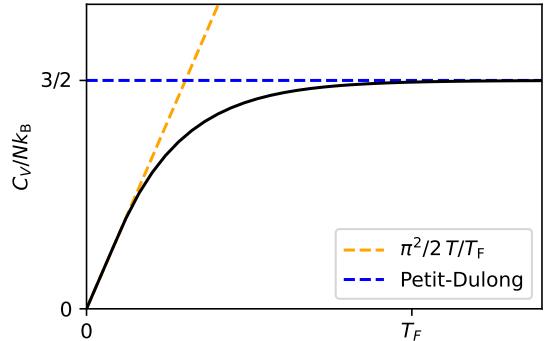
$$\approx \frac{3}{5} \frac{N}{V} E_F \left(1 + \frac{5\pi^2}{12} \left(\frac{k_B T}{E_F} \right)^2 \right) \quad (147)$$

Heat capacity

for low temperatures $T \ll T_F$

$$C_V = \left(\frac{\partial E}{\partial T} \right)_V = N k_B \frac{\pi}{2} \left(\frac{T}{T_F} \right) \quad (148)$$

differs from td:TODO:petit_dulong



TODO:Entartung und Sommerfeld TODO:DULONG-PETIT Gesetz

ed VIII

Electrodynamics

16 Maxwell-Equations

Vacuum
microscopic formulation

$$\vec{\nabla} \cdot \vec{E} = \frac{\rho_{\text{el}}}{\epsilon_0} \quad (149)$$

$$\vec{\nabla} \cdot \vec{B} = 0 \quad (150)$$

$$\vec{\nabla} \times \vec{E} = -\frac{d\vec{B}}{dt} \quad (151)$$

$$\vec{\nabla} \times \vec{B} = \mu_0 \vec{j} + \frac{1}{c^2} \frac{d\vec{E}}{dt} \quad (152)$$

Matter
Macroscopic formulation

$$\vec{\nabla} \cdot \vec{D} = \rho_{\text{el}} \quad (153)$$

$$\vec{\nabla} \cdot \vec{B} = 0 \quad (154)$$

$$\vec{\nabla} \times \vec{E} = -\frac{d\vec{B}}{dt} \quad (155)$$

$$\vec{\nabla} \times \vec{H} = \vec{j} + \frac{d\vec{D}}{dt} \quad (156)$$

17 Fields

17.1 Electric field

Gauss's law for electric fields
Electric flux through a closed surface is proportional to the electric charge

$$\Phi_E = \iint_S \vec{E} \cdot d\vec{S} = \frac{Q}{\epsilon_0} \quad (157)$$

S closed surface

17.2 Electric field

Magnetic flux

$$\Phi_B = \iint_A \vec{B} \cdot d\vec{A} \quad (158)$$

Gauss's law for magnetism
Magnetic flux through a closed surface is 0 \Rightarrow there are no magnetic monopoles

$$\Phi_B = \iint_S \vec{B} \cdot d\vec{S} = 0 \quad (159)$$

S closed surface

(160)

Magnetization

$$\vec{M} = \frac{d\vec{m}}{dV} = \chi_m \cdot \vec{H} \quad (161)$$

m mag. moment, V volume

Torque

$$\vec{\tau} = \vec{m} \times \vec{B} \quad (162)$$

m mag. moment

Susceptibility

$$\chi_m = \frac{\partial M}{\partial B} = \frac{\mu}{\mu_0} - 1 \quad (163)$$

Poynting vector

Directional energy flux or
power flow of an
electromagnetic field [W/m²]

$$\vec{S} = \vec{E} \times \vec{H} \quad (164)$$

17.3 Induction

Faraday's law of induction

$$U_{\text{ind}} = -\frac{d}{dt}\Phi_B = -\frac{d}{dt} \iint_A \vec{B} \cdot d\vec{A} \quad (165)$$

18 Hall-Effect

Cyclotron frequency

$$\omega_c = \frac{eB}{m_e} \quad (166)$$

TODO:Move

18.1 Classical Hall-Effect

Current flowing in x direction in a conductor ($l \times b \times d$) with a magnetic field B in z direction leads to a hall voltage U_H in y direction.

Hall voltage

$$U_H = \frac{IB}{ned} \quad (167)$$

n charge carrier density

Hall coefficient

$$R_H = -\frac{Eg}{j_x B g} = \frac{1}{ne} = \frac{\rho_{xy}}{B_z} \quad (168)$$

Resistivity

$$\rho_{xx} = \frac{m_e}{ne^2 \tau} \quad (169)$$

$$\rho_{xy} = \frac{B}{ne} \quad (170)$$

18.2 Integer quantum hall effect

Conductivity tensor

$$\sigma = \begin{pmatrix} \sigma_{xy} & \sigma_{xy} \\ \sigma_{yx} & \sigma_{yy} \end{pmatrix} \quad (171)$$

Resistivity tensor

$$\rho = \sigma^{-1} \quad (172)$$

Resistivity

$$\rho_{xy} = \frac{2\pi\hbar}{e^2} \frac{1}{\nu} \quad (173)$$

$\nu \in \mathbb{Z}$

TODO:sort

Impedance of a capacitor

$$Z_C = \frac{1}{i\omega C} \quad (174)$$

Impedance of an inductor

$$Z_L = i\omega L \quad (175)$$

TODO:impedance addition for parallel / linear

19 Dipole-stuff

Dipole radiation Poynting vector

$$\vec{S} = \left(\frac{\mu_0 p_0^2 \omega^4}{32\pi^2 c} \right) \frac{\sin^2 \theta}{r^2} \vec{r} \quad (176)$$

Time-average power

$$P = \frac{\mu_0 \omega^4 p_0^2}{12\pi c} \quad (177)$$

qm IX

Quantum Mechanics

20 Basics

20.1 Operators

Dirac notation

$$\langle x| \text{"Bra" Row vector} \quad (178)$$

$$|x\rangle \text{"Ket" Column vector} \quad (179)$$

$$\hat{A}|\beta\rangle = |\alpha\rangle \Rightarrow \langle\alpha| = \langle\beta|\hat{A}^\dagger \quad (180)$$

Dagger

$$\hat{A}^\dagger = (\hat{A}^*)^T \quad (181)$$

$$(c\hat{A})^\dagger = c^* \hat{A}^\dagger \quad (182)$$

$$(\hat{A}\hat{B})^\dagger = \hat{B}^\dagger \hat{A}^\dagger \quad (183)$$

$$(184)$$

Adjoint operator

$$\langle\alpha|\hat{A}^\dagger|\beta\rangle = \langle\beta|\hat{A}|\alpha\rangle^* \quad (185)$$

Hermitian operator

$$\hat{A} = \hat{A}^\dagger \quad (186)$$

20.2 Probability theory

Continuity equation

$$\frac{\partial \rho(\vec{x}, t)}{\partial t} + \nabla \cdot \vec{j}(\vec{x}, t) = 0 \quad (187)$$

ρ density of a conserved quantity q , j flux density of q

State probability

$$TODO \quad (188)$$

Dispersion

$$\Delta\hat{A} = \hat{A} - \langle\hat{A}\rangle \quad (189)$$

Generalized uncertainty principle

$$\sigma_A \sigma_B \geq \frac{1}{4} \langle [\hat{A}, \hat{B}] \rangle^2 \quad (190)$$

$$\sigma_A \sigma_B \geq \frac{1}{2} |\langle [\hat{A}, \hat{B}] \rangle| \quad (191)$$

20.2.1 Pauli matrices

Pauli matrices

$$\sigma_x = \begin{pmatrix} 0 & 1 \\ 1 & 0 \end{pmatrix} = |0\rangle\langle 1| + |1\rangle\langle 0| \quad (192)$$

$$\sigma_y = \begin{pmatrix} 0 & -i \\ i & 0 \end{pmatrix} = -i |0\rangle\langle 1| + i |1\rangle\langle 0| \quad (193)$$

$$\sigma_z = \begin{pmatrix} 1 & 0 \\ 0 & -1 \end{pmatrix} = |0\rangle\langle 0| - |1\rangle\langle 1| \quad (194)$$

20.3 Commutator

Commutator

$$[A, B] = AB - BA \quad (195)$$

Anticommutator

$$\{A, B\} = AB + BA \quad (196)$$

Commutation relations

$$[A, BC] = [A, B]C - B[A, C] \quad (197)$$

TODO: add some more?

Commutator involving a function

$$[f(A), B] = [A, B] \frac{\partial f}{\partial A} \quad (198)$$

given $[A, [A, B]] = 0$

Jacobi identity

$$[A, [B, C]] + [B, [C, A]] + [C, [A, B]] = 0 \quad (199)$$

Hadamard's Lemma

$$e^A B e^{-A} = B + [A, B] + \frac{1}{2!} [A, [A, B]] + \frac{1}{3!} [A, [A, [A, B]]] + \dots \quad (200)$$

Canonical commutation relation

$$[x_i, x_j] = 0 \quad (201)$$

$$[p_i, p_j] = 0 \quad (202)$$

$$[x_i, p_j] = i\hbar\delta_{ij} \quad (203)$$

x, p canonical conjugates

21 Schrödinger equation

Energy operator

$$E = i\hbar \frac{\partial}{\partial t} \quad (204)$$

Momentum operator

$$\vec{p} = -i\hbar \vec{\nabla}_x \quad (205)$$

Space operator

$$\vec{x} = i\hbar \vec{\nabla}_p \quad (206)$$

Stationary Schrödinger equation

$$\hat{H} |\psi\rangle = E |\psi\rangle \quad (207)$$

Schrödinger equation

$$i\hbar \frac{\partial}{\partial t} \psi(x, t) = \left(-\frac{\hbar^2}{2m} \vec{\nabla}^2 + \vec{V}(x)\right) \psi(x) \quad (208)$$

21.1 Time evolution

The time evolution of the Hamiltonian is given by:

Time evolution operator

$$|\psi(t)\rangle = \hat{U}(t, t_0) |\psi(t_0)\rangle \quad (209)$$

U unitary

Von-Neumann Equation
Time evolution of the density operator in the Schrödinger picture. Qm analog to the Liouville equation ??

$$\frac{\partial \hat{\rho}}{\partial t} = -\frac{i}{\hbar} [\hat{H}, \hat{\rho}] \quad (210)$$

Lindblad master equation
Generalization of von-Neumann equation for open quantum systems

$$\dot{\rho} = -\frac{i}{\hbar} [\hat{H}, \rho] + \underbrace{\sum_{n,m} h_{nm} \left(\hat{A}_n \rho \hat{A}_{m^\dagger} - \frac{1}{2} \{ \hat{A}_m^\dagger \hat{A}_n, \rho \} \right)}_{\text{reversible}} \quad (211)$$

h positive semidefinite matrix, \hat{A} arbitrary operator

TODO:unitary transformation of time dependent H

21.1.1 Schrödinger- and Heisenberg-pictures

In the **Schrödinger picture**, the time dependency is in the states while in the **Heisenberg picture** the observables (operators) are time dependent.

Schrödinger time evolution

$$|\psi(t)_S\rangle = \hat{U}(t, t_0) |\psi(t_0)\rangle \quad (212)$$

Heisenberg time evolution

$$|\psi_H\rangle = |\psi_S(t_0)\rangle \quad (213)$$

$$A_H = U^\dagger(t, t_0) A_S U(t, t_0) \quad (214)$$

$$\frac{d\hat{A}_H}{dt} = \frac{1}{i\hbar} [\hat{A}_H, \hat{H}_H] + \left(\frac{\partial \hat{A}_S}{\partial t} \right)_H \quad (215)$$

H and S being the Heisenberg and Schrödinger picture, respectively

21.1.2 Ehrenfest theorem

See also ??

Ehrenfesttheorem applies to both pictures

$$\frac{d}{dt} \langle \hat{A} \rangle = \frac{1}{i\hbar} \langle [\hat{A}, \hat{H}] \rangle + \left\langle \frac{\partial \hat{A}}{\partial t} \right\rangle \quad (216)$$

Example for x

$$m \frac{d^2}{dt^2} \langle x \rangle = -\langle \nabla V(x) \rangle = \langle F(x) \rangle \quad (217)$$

21.2 Correspondence principle

The classical mechanics can be derived from quantum mechanics in the limit of large quantum numbers.

22 Perturbation theory

qm:qm_perturbation:desc

Hamiltonian

$$\hat{H} = \hat{H}_0 + \lambda \hat{H}_1 \quad (218)$$

Power series

$$E_n = E_n^{(0)} + \lambda E_n^{(1)} + \lambda^2 E_n^{(2)} + \dots \quad (219)$$

$$|\psi_n\rangle = |\psi_n^{(0)}\rangle + \lambda |\psi_n^{(1)}\rangle + \lambda^2 |\psi_n^{(2)}\rangle + \dots \quad (220)$$

1. order energy shift

$$E_n^{(1)} = \langle \psi_n^{(0)} | \hat{H}_1 | \psi_n^{(0)} \rangle \quad (221)$$

1. order states

$$|\psi_n^{(1)}\rangle = \sum_{k \neq n} \frac{\langle \psi_k^{(0)} | \hat{H}_1 | \psi_n^{(0)} \rangle}{E_n^{(0)} - E_k^{(0)}} |\psi_k^{(0)}\rangle \quad (222)$$

2. order energy shift

$$E_n^{(2)} = \sum_{k \neq n} \frac{|\langle \psi_k^{(0)} | \hat{H}_1 | \psi_n^{(0)} \rangle|^2}{E_n^{(0)} - E_k^{(0)}} \quad (223)$$

Fermi's golden rule

Transition rate from initial state $|i\rangle$ under a perturbation H^1 to final state $|f\rangle$

$$\Gamma_{i \rightarrow f} = \frac{2\pi}{\hbar} |\langle f | H^1 | i \rangle|^2 \rho(E_f) \quad (224)$$

23 Harmonic oscillator

Hamiltonian

$$H = \frac{p^2}{2m} + \frac{1}{2} m \omega^2 x^2 \quad (225)$$

$$= \frac{1}{2} \hbar \omega + \omega a^\dagger a \quad (226)$$

Energy spectrum

$$E_n = \hbar \omega \left(\frac{1}{2} + n \right) \quad (227)$$

See also ??

23.1 Creation and Annihilation operators / Ladder operators

Particle number
operator/occupation number
operator

$$\hat{N} := a^\dagger a \quad (228)$$

$$\hat{N} |n\rangle = n |N\rangle \quad (229)$$

$|n\rangle$ = Fock states, \hat{a} = Annihilation operator, \hat{a}^\dagger = Creation operator

Commutator

$$[\hat{a}, \hat{a}^\dagger] = 1 \quad (230)$$

$$[N, \hat{a}] = -\hat{a} \quad (231)$$

$$[N, \hat{a}^\dagger] = \hat{a}^\dagger \quad (232)$$

Application on states

$$\hat{a} |n\rangle = \sqrt{n} |n-1\rangle \quad (233)$$

$$\hat{a}^\dagger |n\rangle = \sqrt{n+1} |n+1\rangle \quad (234)$$

$$|n\rangle = \frac{1}{\sqrt{n!}} (\hat{a}^\dagger)^n |0\rangle \quad (235)$$

23.1.1 Harmonischer Oszillatör

Harmonic oscillator

$$\hat{x} = \sqrt{\frac{\hbar}{2m\omega}} (\hat{a} + \hat{a}^\dagger) \quad (236)$$

$$\hat{p} = -i\sqrt{\frac{m\omega\hbar}{2}} (\hat{a} - \hat{a}^\dagger) \quad (237)$$

$$\hat{H} = \frac{\hat{p}^2}{2m} + \frac{m\omega^2 \hat{x}^2}{2} = \hbar\omega \left(a^\dagger a + \frac{1}{2} \right) \quad (238)$$

$$a = \frac{1}{\sqrt{2}} (\tilde{X} + i\tilde{P}) \quad (239)$$

$$a^\dagger = \frac{1}{\sqrt{2}} (\tilde{X} - i\tilde{P}) \quad (240)$$

24 Angular momentum

Bloch waves

Solve the stat. SG in periodic potential with period \vec{R} :

$$V(\vec{r}) = V(\vec{r} + \vec{R})$$

$$\psi_k(\vec{r}) = e^{i\vec{k}\cdot\vec{r}} \cdot u_{\vec{k}}(\vec{r}) \quad (241)$$

\vec{k} arbitrary, u periodic function

24.1 Aharanov-Bohm effect

Acquired phase

Electron along a closed loop
acquires a phase proportional
to the enclosed magnetic flux

$$\delta = \frac{2e}{\hbar} \oint \vec{A} \cdot d\vec{s} = \frac{2e}{\hbar} \Phi \quad (242)$$

TODO: replace with loop integral symbol and add more info

25 Symmetries

Most symmetry operators are unitary ?? because the norm of a state must be invariant under transformations of space, time and spin.

Invariance

\hat{H} is invariant under a
symmetry described by \hat{U} if
this holds

$$\hat{U}\hat{H}\hat{U}^\dagger = \hat{H} \Leftrightarrow [\hat{U}, \hat{H}] = 0 \quad (243)$$

25.1 Time-reversal symmetry

Time-reversal symmetry

$$T : t \rightarrow -t \quad (244)$$

Anti-unitary

$$T^2 = -1 \quad (245)$$

26 Two-level systems (TLS)

James-Cummings

Hamiltonian

TLS interacting with optical
cavity

$$H = \underbrace{\hbar\omega_c \hat{a}^\dagger \hat{a}}_{\text{field}} + \underbrace{\hbar\omega_a \frac{\hat{\sigma}_z}{2}}_{\text{atom}} + \underbrace{\frac{\hbar\Omega}{2} \hat{E} \hat{S}}_{\text{int}} \quad (246)$$

after RWA:

$$(247)$$

$$= \hbar\omega_c \hat{a}^\dagger \hat{a} + \hbar\omega_a \hat{\sigma}^\dagger \hat{\sigma} + \frac{\hbar\Omega}{2} (\hat{a} \hat{\sigma}^\dagger + \hat{a}^\dagger \hat{\sigma}) \quad (248)$$

$\hat{E} = E_{\text{ZPF}}(\hat{a} + \hat{a}^\dagger)$ field operator with bosonic ladder operators,
 $\hat{S} = \hat{\sigma}^\dagger + \hat{\sigma}$ polarization operator with ladder operators
of the TLS

27 Other

Rotating Wave

Approximation (RWS)

Rapidly oscillating terms are neglected

$$\Delta\omega := |\omega_0 - \omega_L| \ll |\omega_0 + \omega_L| \approx 2\omega_0 \quad (249)$$

ω_L light frequency, ω_0 transition frequency

28 Hydrogen Atom

Reduced mass

$$\mu = \frac{m_e m_K}{m_e + m_K} \stackrel{m_e \ll m_K}{\approx} m_e \quad (250)$$

Coulomb potential

For a single electron atom

$$V(\vec{r}) = \frac{Z e^2}{4\pi\epsilon_0 r} \quad (251)$$

Z atomic number

Hamiltonian

$$\hat{H} = -\frac{\hbar^2}{2\mu} \vec{\nabla}_{\vec{r}}^2 - V(\vec{r}) \quad (252)$$

$$= \frac{\hat{p}_r^2}{2\mu} + \frac{\hat{L}^2}{2\mu r} + V(r) \quad (253)$$

Wave function

$$\psi_{nlm}(r, \theta, \phi) = R_{nl}(r) Y_{lm}(\theta, \phi) \quad (254)$$

Radial part

$$R_{nl} = -\sqrt{\frac{(n-l-1)!(2\kappa)^3}{2n[(n+l)!]^3}} (2\kappa r)^l e^{-\kappa r} L_{n+1}^{2l+1}(2\kappa r) \quad (255)$$

with

$$\kappa = \frac{\sqrt{2\mu|E|}}{\hbar} = \frac{Z}{na_B} \quad (256)$$

$L_r^s(x)$ Laguerre-polynomials

Energy eigenvalues

$$E_n = \frac{Z^2 \mu e^4}{n^2 (4\pi\epsilon_0)^2 2\hbar^2} = -E_H \frac{Z^2}{n^2} \quad (257)$$

Rydberg energy

$$E_H = h c R_H = \frac{\mu e^4}{(4\pi\epsilon_0)^2 2\hbar^2} \quad (258)$$

28.1 Corrections

28.1.1 Darwin term

Relativistic correction: Because of the electrons zitterbewegung, it is not entirely localised. **TODO:fact check**

Energy shift

$$\Delta E_{\text{rel}} = -E_n \frac{Z^2 \alpha^2}{n} \left(\frac{3}{4n} - \frac{1}{l + \frac{1}{2}} \right) \quad (259)$$

Fine-structure constant
Sommerfeld constant

$$\alpha = \frac{e^2}{4\pi\epsilon_0 \hbar c} \approx \frac{1}{137} \quad (260)$$

28.1.2 Spin-orbit coupling (LS-coupling)

The interaction of the electron spin with the electrostatic field of the nuclei lead to energy shifts.

Energy shift

$$\Delta E_{\text{LS}} = \frac{\mu_0 Z e^2}{8\pi m_e^2 r^3} \langle \vec{S} \cdot \vec{L} \rangle \quad (261)$$

TODO:name

$$\begin{aligned} \langle \vec{S} \cdot \vec{L} \rangle &= \frac{1}{2} \langle [J^2 - L^2 - S^2] \rangle \\ &= \frac{\hbar^2}{2} [j(j+1) - l(l+1) - s(s+1)] \end{aligned} \quad (262)$$

28.1.3 Fine-structure

The fine-structure combines relativistic corrections 28.1.1 and the spin-orbit coupling 28.1.2.

Energy shift

$$\Delta E_{\text{FS}} = \frac{Z^2 \alpha^2}{n} \left(\frac{1}{j + \frac{1}{2}} - \frac{3}{4n} \right) \quad (263)$$

28.1.4 Lamb-shift

The interaction of the electron with virtual photons emitted/absorbed by the nucleus leads to a (very small) shift in the energy level.

Potential energy

$$\langle E_{\text{pot}} \rangle = -\frac{Ze^2}{4\pi\epsilon_0} \left(\frac{1}{r + \delta r} \right) \quad (264)$$

δr perturbation of r

28.1.5 Hyperfine structure

Interaction of the nucleus spin with the magnetic field created by the electron leads to energy shifts.
(Lifts degeneracy)

Nuclear spin

$$\vec{F} = \vec{J} + \vec{I} \quad (265)$$

$$|\vec{I}| = \sqrt{i(i+1)}\hbar \quad (266)$$

$$I_z = m_i \hbar \quad (267)$$

$$m_i = -i, -i+1, \dots, i-1, i \quad (268)$$

Combined angular momentum

$$\vec{F} = \vec{J} + \vec{I} \quad (269)$$

$$|\vec{F}| = \sqrt{f(f+1)}\hbar \quad (270)$$

$$F_z = m_f \hbar \quad (271)$$

Selection rule

$$f = j \pm i \quad (272)$$

$$m_f = -f, -f+1, \dots, f-1, f \quad (273)$$

Hyperfine structure constant

$$A = \frac{g_i \mu_K B_{\text{HFS}}}{\sqrt{j(j+1)}} \quad (274)$$

B_{HFS} hyperfine field, μ_K nuclear magneton, g_i nuclear g-factor ??

Energy shift

$$\Delta H_{\text{HFS}} = \frac{A}{2} [f(f+1) - j(j+1) - i(i+1)] \quad (275)$$

TODO:landé factor

28.2 Effects in magnetic field

TODO:all

TODO:Hunds rules

Condensed matter physics

29 Bravais lattice

Table 2: In 2D, there are 5 different Bravais lattices

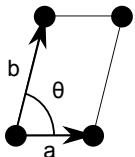
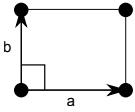
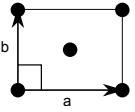
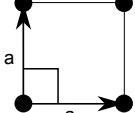
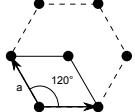
Lattice system	Point group	5 Bravais lattices	
		primitive (p)	centered (c)
monoclinic (m)	C_2		
orthorhombic (o)	D_2		
tetragonal (t)	D_4		
hexagonal (h)	D_6		

Table 3: In 3D, there are 14 different Bravais lattices

Crystal system	Lattice system	Point group	14 Bravais lattices			
			primitive (P)	base_centered (S)	body_centered (I)	face_centered (F)
triclinic (a)		C _i				
monoclinic (m)		C _{2h}				
orthorhombic (o)		D _{2h}				
tetragonal (t)		D _{4h}				
hexagonal (h)	rhombohedral	D _{3d}				
	hexagonal	D _{6h}				
cubic (c)		O _h				

30 Reciprocal lattice

The reciprocal lattice is made up of all the wave vectors \vec{k} that resemble standing waves with the periodicity of the Bravais lattice.

Reciprocal lattice vectors

$$\vec{b}_1 = \frac{2\pi}{V_c} \vec{a}_2 \times \vec{a}_3 \quad (276)$$

$$\vec{b}_2 = \frac{2\pi}{V_c} \vec{a}_3 \times \vec{a}_1 \quad (277)$$

$$\vec{b}_3 = \frac{2\pi}{V_c} \vec{a}_1 \times \vec{a}_2 \quad (278)$$

a_i real-space lattice vectors, V_c volume of the primitive lattice cell

30.1 Scattering processes

Matthiessen's rule

Approximation, only holds if the processes are independent of each other

$$\frac{1}{\mu} = \sum_{i=\text{Scattering processes}} \frac{1}{\mu_i} \quad (279)$$

$$\frac{1}{\tau} = \sum_{i=\text{Scattering processes}} \frac{1}{\tau_i} \quad (280)$$

μ mobility, τ scattering time

31 Free electron gas

Assumptions: electrons can move freely and independent of each other.

Drift velocity

Velocity component induced by an external force (eg. electric field)

$$\vec{v}_D = \vec{v} - \vec{v}_{th} \quad (281)$$

v_{th} thermal velocity

Mean free time

$$\tau \quad (282)$$

Mean free path

$$\ell = \langle v \rangle \tau \quad (283)$$

Electrical mobility

$$\mu = \frac{q\tau}{m} \quad (284)$$

q charge, m mass

31.1 Drude model

Classical model describing the transport properties of electrons in materials (metals): The material is assumed to be an ion lattice and with freely moving electrons (electron gas). The electrons are accelerated by an electric field and decelerated through collisions with the lattice ions. The model disregards the Fermi-Dirac partition of the conducting electrons.

Equation of motion

$$m_e \frac{d\vec{v}}{dt} + \frac{m_e}{\tau} \vec{v}_D = -e \vec{E} \quad (285)$$

v electron speed, \vec{v}_D drift velocity, τ mean free time between collisions

Current density

Ohm's law

$$\vec{j} = -ne\vec{v}_D = ne\mu\vec{E} \quad (286)$$

n charge particle density

Drude-conductivity

$$\sigma = \frac{\vec{j}}{\vec{E}} = \frac{e^2 \tau n}{m_e} = ne\mu \quad (287)$$

31.2 Sommerfeld model

Assumes a gas of free fermions underlying the pauli-exclusion principle. Only electrons in an energy range of $k_B T$ around the Fermi energy E_F participate in scattering processes.

Current density

$$\vec{j} = -en \langle v \rangle = -en \frac{\hbar}{m_e} \langle \vec{k} \rangle = -e \frac{1}{V} \sum_{\vec{k}, \sigma} \frac{\hbar \vec{k}}{m_e} \quad (288)$$

TODO: The formula for the conductivity is the same as in the drude model?

31.3 2D electron gas

Lower dimension gases can be obtained by restricting a 3D gas with infinitely high potential walls on a narrow area with the width L .

Confinement energy
Raises ground state energy

$$\Delta E = \frac{\hbar^2 \pi^2}{2m_e L^2} \quad (289)$$

Energy

$$E_n = \underbrace{\frac{\hbar^2 k_{\parallel}^2}{2m_e}}_{x-y: \text{ plain wave}} + \underbrace{\frac{\hbar^2 \pi^2}{2m_e L^2} n^2}_{z} \quad (290)$$

31.4 1D electron gas / quantum wire

Energy

$$E_n = \frac{\hbar^2 k_x^2}{2m_e} + \frac{\hbar^2 \pi^2}{2m_e L_z^2} n_1^2 + \frac{\hbar^2 \pi^2}{2m_e L_y^2} n_2^2 \quad (291)$$

31.5 0D electron gas / quantum dot

TODO:TODO

32 Measurement techniques

32.1 ARPES

what? in? how? plot

32.2 Scanning probe microscopy SPM

Images of surfaces are taken by scanning the specimen with a physical probe.

Name	Atomic force microscopy (AFM)
Application	Surface stuff
how	With needle

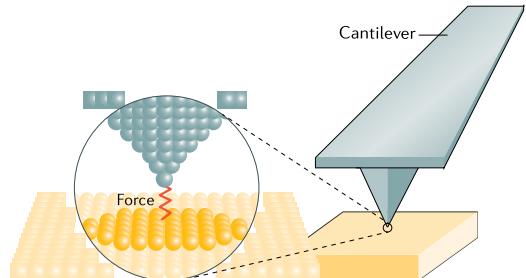


Figure 1: [?]

Name	Scanning tunneling microscopy (STM)
Application	Surface stuff
how	With Tunnel

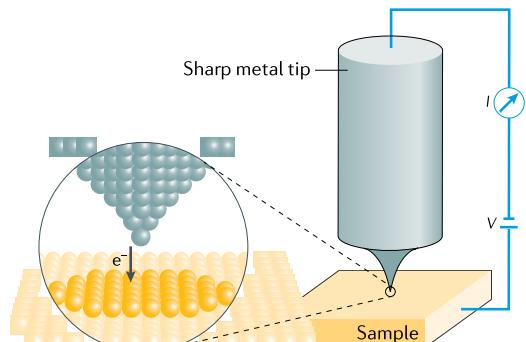
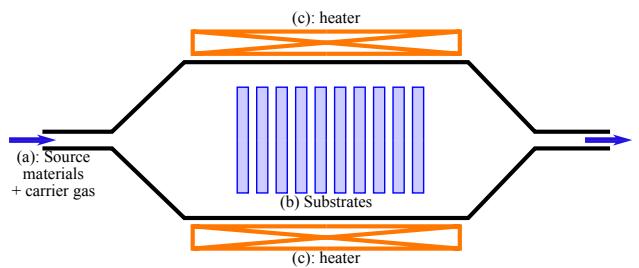


Figure 2: [?]

33 Fabrication techniques

Name	Chemical vapor deposition (CVD)
how	A substrate is exposed to volatile precursors, which react and/or decompose on the heated substrate surface to produce the desired deposit. By-products are removed by gas flow through the chamber.
Application	<ul style="list-style-type: none"> • Polysilicon Si • Silicon dioxide SiO_2 • Graphene • Diamond



33.1 Epitaxy

A type of crystal growth in which new layers are formed with well-defined orientations with respect to the crystalline seed layer.

Name	Molecular Beam Epitaxy (MBE)
how	In a ultra-high vacuum, the elements are heated until they slowly sublime. The gases then condensate on the substrate surface
Application	<ul style="list-style-type: none"> • Gallium arsenide GaAs <p style="color: red;">TODO:Link to GaAs</p>

