

# Thermal field theory, WS 08/09

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## I. INTRODUCTION AND OUTLINE

We shall discuss systems in equilibrium at finite temperatures and chemical potentials. In most parts we will follow the book by Kapusta [1], see also the book by Le Bellac [2] for additional reference. We will focus on the functional integral approach, for a different approach using second quantization see the book by Fetter and Walecka [3]. We

shall learn the tools of functional integration, Matsubara summation, perturbation techniques, and discuss important theoretical concepts such as spontaneous symmetry breaking (and restoration thereof at large temperatures). Applications, to be discussed after learning these techniques, are

- early universe, cosmology
  - inflation,  $T \sim 10^{15}$  GeV
  - electroweak phase transition  $T \sim 10^2$  GeV
  - QCD phase transition,  $T \sim 10^2$  MeV (for comparison, this is  $\sim 10^{12}$  K)
  - baryogenesis
- QCD phase transitions
  - heavy-ion collisions (“little bang” vs. “big bang”)
  - chiral symmetry (spontaneous breaking thereof)
  - lattice QCD
- neutron stars,  $T \lesssim 10$  MeV,  $\mu_q \sim 400$  MeV
  - dense nuclear matter
  - neutrino emissivity
  - quark matter, color superconductivity

In this lecture we can only touch a few of these points.

## II. REMINDER: BASICS OF STATISTICAL QUANTUM MECHANICS

This chapter serves as a quick reminder of the main ingredients of statistical quantum mechanics and its relation to thermodynamic quantities. A good textbook about statistical physics is the book by Nolting [4]. The goal of this reminder is to explain the meaning and form of the partition function which, in later chapters in the functional integral representation, plays a central role in thermal field theory.

### A. Statistical operator

We start by recalling that statistical quantum mechanics involves probabilities on two levels. First, on the fundamental level, quantum mechanics itself involves some kind of statistics, i.e., we can only give probabilities to measure a certain value of an observable. Let  $\hat{A}$  be an observable with a set of complete orthogonal eigenstates  $|n\rangle$  and eigenvalues  $a_n$ ,

$$\hat{A}|n\rangle = a_n|n\rangle, \quad (1)$$

with  $\langle m|n\rangle = \delta_{mn}$ . We can expand any state  $|\psi\rangle$  in terms of these eigenstates,

$$|\psi\rangle = \sum_n c_n |n\rangle. \quad (2)$$

The probability to measure the value  $a_n$  in the state  $|\psi\rangle$  is  $|c_n|^2$  with  $c_n = \langle n|\psi\rangle$ . Then, the expectation value of  $\hat{A}$  in the state  $|\psi\rangle$  is

$$\begin{aligned} \langle \hat{A} \rangle &= \sum_n |c_n|^2 a_n \\ &= \sum_n |\langle n|\psi\rangle|^2 a_n \\ &= \sum_n \langle n|\psi\rangle \langle \psi|\hat{A}|n\rangle \\ &= \langle \psi|\hat{A}|\psi\rangle. \end{aligned} \quad (3)$$

Second, and independent of the quantum probability interpretation, there is a level of uncertainty for macroscopic systems of which we don't know (and are not interested in) the microscopic details. We now consider many possible (orthogonal) quantum mechanical states  $|\psi_m\rangle$  each of which we find with a probability  $p_m$ ,  $0 < p_m < 1$ . Then, the expectation value for  $\hat{A}$  is not only given by the quantum mechanical averaging but also by averaging over the possible states  $|\psi_m\rangle$ , i.e.,

$$\langle \hat{A} \rangle = \sum_m p_m \langle \psi_m | \hat{A} | \psi_m \rangle. \quad (4)$$

If we define the statistical operator

$$\hat{\rho} \equiv \sum_m p_m |\psi_m\rangle \langle \psi_m|, \quad (5)$$

we can write the expectation value as

$$\langle \hat{A} \rangle = \text{Tr}(\hat{\rho} \hat{A}). \quad (6)$$

Proof:

$$\begin{aligned} \langle \hat{A} \rangle &= \sum_m p_m \langle \psi_m | \hat{A} | \psi_m \rangle = \sum_{i,j,m} p_m \langle \psi_m | \phi_i \rangle \langle \phi_i | \hat{A} | \phi_j \rangle \langle \phi_j | \psi_m \rangle \\ &= \sum_{i,j} \hat{\rho}_{ji} \hat{A}_{ij} = \text{Tr}(\hat{\rho} \hat{A}). \end{aligned} \quad (7)$$

We obviously have  $\hat{\rho}^\dagger = \hat{\rho}$  and

$$\text{Tr} \hat{\rho} = 1 \quad (8)$$

(which is clear from setting  $\hat{A} = \mathbf{1}$  above). The meaning of  $\hat{\rho}$  can also be understood from the analogy to classical statistical mechanics. In this case, there is a probability distribution  $\rho(p, q)$  in the  $6N$ -dimensional phase space ( $N$  be the number of particles which each moves on a trajectory in phase space); then,  $d^{3N}p d^{3N}q \rho(p, q)$  is the probability to find the system in the small region  $d^{3N}p d^{3N}q$  in phase space. An observable  $A$  has a value  $A(p, q)$  if the system sits on the point  $(p, q)$  in phase space, and its expectation value is given by

$$\langle A \rangle = \frac{1}{(2\pi\hbar)^{3N} N!} \int d^{3N}p d^{3N}q \rho(p, q) A(p, q), \quad (9)$$

where the factor  $N!$  in the normalization refers to the exchange of particles and the factor  $(2\pi\hbar)^{3N}$  is included to do the transition to a quantum mechanical system. Comparing Eq. (6) with Eq. (9) shows the formal similarity between the quantum and classical expectation values (the trace is replaced by the phase space integral). This formal correspondence goes further, e.g., the Liouville equation for the classical probability distribution

$$\frac{\partial \rho}{\partial t} = -\{H, \rho\}, \quad (10)$$

where  $H$  is the Hamilton function of the system and  $\{-, -\}$  the Poisson bracket, is very similar to the Heisenberg equation for the statistical operator

$$\frac{\partial \hat{\rho}}{\partial t} = -\frac{i}{\hbar} [\hat{H}, \hat{\rho}], \quad (11)$$

with the Hamilton operator  $\hat{H}$ . Here we shall be mostly concerned with equilibrium situations and thus  $[\hat{H}, \hat{\rho}] = 0$ .

## B. Grand canonical ensemble

Let us now recall the different ensembles in statistical physics (both classical and quantum mechanical). One should in the following always think of an ensemble as a collection of many different systems with the same fixed macroscopic properties but different microscopic configurations. Which macroscopic properties are fixed depends on the ensemble:

- microcanonical ensemble  $(E, N, V)$
- canonical ensemble  $(T, N, V)$
- grand canonical ensemble  $(T, \mu, V)$

with the volume  $V$ , the energy  $E$ , the particle number (charge)  $N$ , the temperature  $T$ , and the chemical potential  $\mu$  (associated with the charge  $N$ ); in general, there can be more than one conserved charge and chemical potential. We shall mostly be concerned with the grand canonical ensemble. Therefore, let us give a brief derivation of the statistical operator in the grand canonical ensemble. Consider a system  $\Sigma$  with fixed energy  $E$ , charge  $N$ , and volume  $V$ . We are interested in a small subsystem  $\Sigma^{(1)}$  such that  $\Sigma = \Sigma^{(1)} \cup \Sigma^{(2)}$  and such that  $\Sigma^{(1)}$  and  $\Sigma^{(2)}$  are separated by walls through which charge and energy can be exchanged. Let us denote the energy, charge, and volume of the subsystems by  $E^{(i)}$ ,  $N^{(i)}$ ,  $V^{(i)}$ . We then have  $E = E^{(1)} + E^{(2)}$ ,  $N = N^{(1)} + N^{(2)}$ . We assume that the Hamilton and charge operators commute,  $[\hat{H}^{(i)}, \hat{N}^{(i)}] = 0$ , and thus that there is a set of simultaneous eigenstates  $|E_m, n\rangle$  for the subsystem  $\Sigma_1$ ,

$$\hat{H}^{(1)}|E_m, n\rangle = E_m|E_m, n\rangle, \quad (12a)$$

$$\hat{N}^{(1)}|E_m, n\rangle = n|E_m, n\rangle. \quad (12b)$$

We can write the statistical operator as

$$\hat{\rho} = \sum_{m,n} p_{m,n} |E_m, n\rangle \langle E_m, n|. \quad (13)$$

We are interested in the probability  $p_{m,n}$  to find the system  $\Sigma^{(1)}$  in the state  $|E_m, n\rangle$  with energy  $E_m$  and charge  $N^{(1)} = n$ . This probability is proportional to the number of states  $\Gamma$  available in the complementary system  $\Sigma^{(2)}$ ,

$$p_{m,n} \propto \Gamma^{(2)}(E - E_m, N - n, V^{(2)}). \quad (14)$$

The system  $\Sigma^{(2)}$  acts as a heat and particle bath for  $\Sigma^{(1)}$  and thus  $E_m \ll E$ ,  $n \ll N$ . We can then expand the logarithm of  $\Gamma$ ,

$$\ln \Gamma^{(2)}(E - E_m, N - n, V^{(2)}) \simeq \frac{1}{k_B} S^{(2)}(E, N, V^{(2)}) - \frac{E_m}{k_B} \left. \frac{\partial S^{(2)}}{\partial E} \right|_{E_m=n=0} - \frac{n}{k_B} \left. \frac{\partial S^{(2)}}{\partial N} \right|_{E_m=n=0}, \quad (15)$$

where we have used the definition for the entropy  $S = k_B \ln \Gamma$  where  $k_B$  is the Boltzmann constant. Now we can use the thermodynamic relations  $\partial S / \partial E = 1/T$ ,  $\partial S / \partial N = -\mu/T$  (or one can even view the derivatives in Eq. (15) as the definitions of temperature and chemical potential, which might be a bit abstract) to obtain

$$p_{m,n} \propto e^{-\beta(E_m - \mu n)}. \quad (16)$$

Here we have dropped the contribution  $S^{(2)}/k_B$  which only depends on the system  $\Sigma^{(2)}$  and thus can be absorbed in the normalization, to be determined below. Inserting this into the statistical operator (13) yields

$$\hat{\rho} \propto \sum_{m,n} e^{-\beta(E_m - \mu n)} |E_m, n\rangle \langle E_m, n| = e^{-\beta(\hat{H}^{(1)} - \mu \hat{N}^{(1)})} \sum_{m,n} |E_m, n\rangle \langle E_m, n| = e^{-\beta(\hat{H}^{(1)} - \mu \hat{N}^{(1)})}. \quad (17)$$

To fulfill the normalization (8) we find (dropping the superscript (1), from now on we are only talking about the subsystem  $\Sigma^{(1)}$ )

$$\hat{\rho} = \frac{e^{-\beta(\hat{H} - \mu \hat{N})}}{Z}, \quad (18)$$

where

$$Z \equiv \text{Tr} e^{-\beta(\hat{H} - \mu \hat{N})} \quad (19)$$

is the partition function of the system.

[End of 1st lecture, Oct 6th, 2008.]

We can derive all thermodynamical quantities from the partition function, which thus is the central quantity in statistical physics. For instance we have the grand canonical potential (sometimes called the thermodynamical potential)

$$\Omega(T, \mu, V) \equiv -\frac{1}{\beta} \ln Z. \quad (20)$$

This function is related to the other thermodynamical quantities via

$$\Omega = -PV = E - \mu N - TS, \quad (21)$$

where  $P$  is the pressure. From the partition function we immediately get, in accordance with (21),

$$\begin{aligned} \frac{\partial \Omega}{\partial \mu} &= -\frac{1}{\beta} \frac{\partial \ln Z}{\partial \mu} \\ &= -\frac{1}{\beta Z} \frac{\partial Z}{\partial \mu} \\ &= -\frac{\text{Tr} \hat{N} e^{-\beta(\hat{H} - \mu \hat{N})}}{Z} \\ &= -\langle \hat{N} \rangle \equiv -N, \end{aligned} \quad (22)$$

and

$$\frac{\partial \Omega}{\partial T} = -\ln Z - \beta \frac{\text{Tr} \left[ (\hat{H} - \mu \hat{N}) e^{-\beta(\hat{H} - \mu \hat{N})} \right]}{Z} = \langle \ln \hat{\rho} \rangle = -\langle S \rangle, \quad (23)$$

since  $\ln \hat{\rho} = \ln e^{-\beta(\hat{H} - \mu \hat{N})} - \ln Z = -\beta(\hat{H} - \mu \hat{N}) - \ln Z$ .

Let us compute the partition function for the simplest cases before we turn to the field theoretical description. Let us first consider a single energy state with energy  $\omega$  which we can fill with bosons. Remember that this simple many-particle system resembles the one-particle harmonic oscillator because the Hamiltonian is  $\hat{H} = \omega(\hat{N} + 1/2)$ , where we drop the zero-point energy  $\omega/2$ . Since arbitrarily many bosons can populate the state the partition function is

$$Z = \sum_{n=0}^{\infty} \langle n | e^{-\beta(\omega - \mu)\hat{N}} | n \rangle = \sum_{n=0}^{\infty} e^{-\beta(\omega - \mu)n} \langle n | n \rangle = \frac{1}{1 - e^{-\beta(\omega - \mu)}}. \quad (24)$$

The thermodynamical potential is thus

$$\Omega = -T \ln Z = T \ln \left[ 1 - e^{-\beta(\omega - \mu)} \right] \quad (25)$$

and the particle number

$$N = -\frac{\partial \Omega}{\partial \mu} = \frac{1}{e^{\beta(\omega - \mu)} - 1}. \quad (26)$$

One can already see from this simple example that something interesting happens if the chemical potential approaches the energy  $\omega$ , since in this case the particle number seems to diverge. This is a first hint of Bose-Einstein condensation which we shall discuss later. We also see that the chemical potential cannot assume values larger than  $\omega$  in order to avoid negative  $N$ . This is a restriction for non-interacting systems.

For fermions we have the same expression, only that we can only put one fermion at most into the energy state. Consequently, the sum only runs over  $n = 0, 1$  and we get

$$Z = \sum_{n=0}^1 \langle n | e^{-\beta(\omega - \mu)\hat{N}} | n \rangle = 1 + e^{-\beta(\omega - \mu)}, \quad (27)$$

where the  $n = 0$  term yields the 1 from the expansion of the exponential (which is obvious from writing  $e^{-\beta(\omega - \mu)\hat{N}} = \mathbf{1} - \beta(\omega - \mu)\hat{N} + \dots$ ). In this case, the thermodynamical potential is

$$\Omega = -T \ln Z = -T \ln \left[ 1 + e^{-\beta(\omega - \mu)} \right] \quad (28)$$

and the particle number

$$N = -\frac{\partial \Omega}{\partial \mu} = \frac{1}{e^{\beta(\omega-\mu)} + 1}. \quad (29)$$

In this simple case of just one single energy  $\omega$ , the particle number at vanishing temperature is  $N = 0$  for  $\mu < \omega$  and  $N = 1$  for  $\mu > \omega$ . At nonzero temperature,  $N$  assumes values in between 0 and 1.

Next we allow for a dispersion of the particles, i.e., the energy may depend on the (modulus of the) momentum,  $\omega_p \equiv \omega(p)$ . We start with a box with size  $L$  in all three dimensions. The box size must be an integer times half of the wavelength  $\lambda$ ,  $L = n\lambda/2$ . With the de Broglie relation for the wavelength  $p = 2\pi/\lambda$  (we set  $\hbar = 1$ ) we have  $p = n\pi/L$ . The log of the full partition function is the sum over all partition functions of the single modes,

$$\ln Z = \sum_{\mathbf{n}=(n_x, n_y, n_z)} \ln Z_{\mathbf{n}} = V \int \frac{d^3 p}{(2\pi)^3} \ln Z_p = \mp V \sum_{e=\pm} \int \frac{d^3 p}{(2\pi)^3} \ln[1 \mp e^{-\beta(\omega_p - e\mu)}], \quad (30)$$

where  $V = L^3$ , where we have used " $dn \rightarrow L/\pi dp$ ", and where the doubling of all three momentum ranges in the integral yields the factor  $2^3$  in the denominator. We have inserted the above expressions for the log of the partition function for a single mode for bosons (upper sign) and fermions (lower sign). Finally, we have added a sum over  $e = \pm$  accounting for particles and antiparticles which differ in the sign of their chemical potential. The conserved charge is thus

$$N = V \sum_{e=\pm} e \int \frac{d^3 p}{(2\pi)^3} \frac{1}{e^{\beta(\omega_p - e\mu)} \mp 1}. \quad (31)$$

We shall often consider the charge density  $n \equiv N/V$  instead. In the case of bosons (upper sign) we see that we have to require  $-\min \omega_p < \mu < \min \omega_p$  in order to have positive occupation numbers.

### III. PARTITION FUNCTION AS FUNCTIONAL INTEGRAL

Here we derive the expression for the partition function in quantum field theory as opposed to usual quantum mechanics. Remember that in usual quantum theory, the projection of an eigenstate  $\langle \mathbf{x} |$  of the position operator  $\hat{\mathbf{x}}$  onto the eigenstate  $|\mathbf{p}\rangle$  of the momentum operator  $\hat{\mathbf{p}}$  is given by a plane wave,

$$\langle \mathbf{x} | \mathbf{p} \rangle = e^{i\mathbf{p} \cdot \mathbf{x}}. \quad (32)$$

In quantum field theory, the discrete sum  $\mathbf{p} \cdot \mathbf{x} = \sum_i p_i x_i$  becomes an integral,

$$\langle \phi | \pi \rangle = e^{i \int d^3 x \pi(\mathbf{x}) \phi(\mathbf{x})}. \quad (33)$$

Here,  $\phi(\mathbf{x})$  and  $\pi(\mathbf{x})$  are eigenvalues (better: eigenfunctions) of the field operator (at  $t = 0$ )  $\hat{\phi}(\mathbf{x}, 0)$  and its conjugate momentum operator  $\hat{\pi}(\mathbf{x}, 0)$ . For the completeness and orthogonality conditions one needs functional integration,

$$\int d\phi(\mathbf{x}) |\phi\rangle \langle \phi| = 1, \quad \langle \phi_a | \phi_b \rangle = \delta[\phi_a(\mathbf{x}) - \phi_b(\mathbf{x})], \quad (34a)$$

$$\int \frac{d\pi(\mathbf{x})}{2\pi} |\pi\rangle \langle \pi| = 1, \quad \langle \pi_a | \pi_b \rangle = \delta[\pi_a(\mathbf{x}) - \pi_b(\mathbf{x})]. \quad (34b)$$

The Hamiltonian  $\hat{H}$  of the system is given by the Hamilton density  $\mathcal{H}$  which can be expressed in terms of the field operators,

$$\hat{H} = \int d^3 x \mathcal{H}(\hat{\pi}(\mathbf{x}, t), \hat{\phi}(\mathbf{x}, t)). \quad (35)$$

In the following we shall write the partition function in terms of the fields  $\phi$  and  $\pi$  and get rid of all operators. To this end, we first compute a transition amplitude with identical initial and final state  $\phi$  at times  $t = 0$  and  $t = t_f$ . The initial state evolves in time upon applying the unitary operator  $e^{-i\hat{H}t}$ . We shall divide the time interval  $[0, t_f]$

into  $N$  pieces with length  $\Delta t$ . Then we can write the transition amplitude as

$$\begin{aligned} \langle \phi | e^{-i\hat{H}t} | \phi \rangle &= \lim_{N \rightarrow \infty} \langle \phi | e^{-i\hat{H}\Delta t} e^{-i\hat{H}\Delta t} \dots e^{-i\hat{H}\Delta t} | \phi \rangle \\ &= \int \prod_{i=1}^N \frac{d\pi_i(\mathbf{x})}{2\pi} d\phi_i(\mathbf{x}) \langle \phi | \pi_N \rangle \langle \pi_N | e^{-i\hat{H}\Delta t} | \phi_N \rangle \langle \phi_N | \pi_{N-1} \rangle \langle \pi_{N-1} | e^{-i\hat{H}\Delta t} | \phi_{N-1} \rangle \\ &\quad \times \dots \times \langle \phi_2 | \pi_1 \rangle \langle \pi_1 | e^{-i\hat{H}\Delta t} | \phi_1 \rangle \langle \phi_1 | \phi \rangle, \end{aligned} \quad (36)$$

where we inserted each of the completeness relations in Eqs. (34)  $N$  times alternately. Now, for the scalar products of the form  $\langle \phi_{i+1} | \pi_i \rangle$  we use Eq. (33). For the factors involving the Hamiltonian we use

$$\begin{aligned} \langle \pi_i | e^{-i\hat{H}\Delta t} | \phi_i \rangle &= e^{-i\Delta t \int d^3x \mathcal{H}(\phi_i, \pi_i)} \langle \pi_i | \phi_i \rangle \\ &= e^{-i\Delta t \int d^3x \mathcal{H}(\phi_i, \pi_i)} e^{i \int d^3x \pi_i \phi_i} \end{aligned} \quad (37)$$

Finally from the last product in the integrand in Eq. (36) we obtain  $\langle \phi_1 | \phi \rangle = \delta(\phi - \phi_1)$ . Consequently,

$$\begin{aligned} \langle \phi | e^{-i\hat{H}t} | \phi \rangle &= \lim_{N \rightarrow \infty} \int \prod_{i=1}^N \frac{d\pi_i(\mathbf{x})}{2\pi} d\phi_i(\mathbf{x}) \delta[\phi(\mathbf{x}) - \phi_1(\mathbf{x})] e^{i \int d^3x [\pi_N(\phi - \phi_N) + \pi_{N-1}(\phi_N - \phi_{N-1}) + \dots + \pi_1(\phi_2 - \phi_1)]} \\ &\quad \times e^{-i\Delta t \int d^3x [\mathcal{H}(\phi_N, \pi_N) + \dots + \mathcal{H}(\phi_1, \pi_1)]} \\ &= \lim_{N \rightarrow \infty} \int \prod_{i=1}^N \frac{d\pi_i(\mathbf{x})}{2\pi} d\phi_i(\mathbf{x}) \delta[\phi(\mathbf{x}) - \phi_1(\mathbf{x})] \exp \left\{ i\Delta t \sum_{j=1}^N \int d^3x \left[ \pi_j \frac{\phi_{j+1} - \phi_j}{\Delta t} - \mathcal{H}(\phi_j, \pi_j) \right] \right\}, \end{aligned} \quad (38)$$

where we have denoted  $\phi_{N+1} \equiv \phi$ . We can now take the limit  $N \rightarrow \infty$  to obtain

$$\langle \phi | e^{-i\hat{H}t} | \phi \rangle = \int \mathcal{D}\pi \int_{\phi(\mathbf{x},0)=\phi(\mathbf{x})}^{\phi(\mathbf{x},t_f)=\phi(\mathbf{x})} \mathcal{D}\phi \exp \left\{ i \int_0^{t_f} dt \int d^3x [\pi(\mathbf{x},t) \partial_t \phi(\mathbf{x},t) - \mathcal{H}(\phi(\mathbf{x},t), \pi(\mathbf{x},t))] \right\}. \quad (39)$$

We have denoted the continuum limit of the functional integration as

$$\int \prod_{i=1}^N \frac{d\pi_i(\mathbf{x})}{2\pi} \rightarrow \mathcal{D}\pi, \quad \int \prod_{i=1}^N d\phi_i(\mathbf{x}) \rightarrow \mathcal{D}\phi. \quad (40)$$

From this definition of the continuum limit we also learn, as an aside, how functional integration is defined in general. Note that in the current derivation we have discretized the time direction since this is the crucial direction in order to compute the partition function at finite  $T$ , see below. The functional integration with respect to the space direction is defined analogously, i.e., by taking the continuum limit from discretized space.

We can now use the result (39) to compute the partition function. To this end we compare Eq. (39) with Eq. (19). We see that the trace looks like a transition amplitude with identical initial and final states,

$$\begin{aligned} Z &= \text{Tr} e^{-\beta(\hat{H} - \mu\hat{N})} \\ &= \int d\phi \langle \phi | e^{-\beta(\hat{H} - \mu\hat{N})} | \phi \rangle \\ &= \int \mathcal{D}\pi \int_{\text{periodic}} \mathcal{D}\phi \exp \left[ - \int_0^\beta d\tau \int d^3x (\mathcal{H} - \mu\mathcal{N} - i\pi \partial_\tau \phi) \right]. \end{aligned} \quad (41)$$

Here, we have, upon comparing with Eq. (39), identified the inverse temperature with ‘‘imaginary time’’  $\tau = it$ , such that the integration over  $\tau$  goes from 0 to the inverse temperature  $\beta = 1/T$ . The term ‘‘periodic’’ for the  $\phi$  integral means that all functions  $\phi$  have to be periodic in the imaginary time direction,  $\phi(\mathbf{x}, 0) = \phi(\mathbf{x}, \beta)$ . The integral over  $d\phi$  integrates over all boundary values which are fixed in Eq. (39). We are left with a partition function which is given entirely in terms of the fields, all operators are gone.

#### IV. REAL NON-INTERACTING SCALAR FIELD

We now compute the partition function (41) for the simplest case, a real non-interacting scalar field which is described by the Lagrangian

$$\mathcal{L} = \frac{1}{2} \partial_\mu \phi \partial^\mu \phi - \frac{1}{2} m^2 \phi^2 = \frac{1}{2} [(\partial_0 \phi)^2 - (\nabla \phi)^2 - m^2 \phi^2]. \quad (42)$$

In this simple case there is no conserved charge, hence there is no chemical potential. We shall introduce the chemical potential for a charged complex field in the subsequent section. The conjugate momentum is

$$\pi = \frac{\partial \mathcal{L}}{\partial(\partial_0 \phi)} = \partial_0 \phi. \quad (43)$$

Remember that  $\mathcal{L}$  and  $\mathcal{H}$  are connected via a Legendre transformation which changes the independent variable  $\partial_0 \phi$  (velocity  $\dot{q}$  in classical mechanics) to  $\pi$  (momentum  $p$  in classical mechanics). Therefore, the Hamiltonian is

$$\mathcal{H} = \pi \partial_0 \phi - \mathcal{L} = \frac{1}{2} [\pi^2 + (\nabla \phi)^2 + m^2 \phi^2]. \quad (44)$$

For the partition function (41) we need the combination  $\pi \partial_0 \phi - \mathcal{H}$ . This looks like the Lagrangian, but the variables over which we integrate are  $\pi$  and  $\phi$ , i.e., we have to be careful with identifying the Lagrangian in the integrand. In order to do so, we can shift the momentum field by introducing

$$\tilde{\pi} \equiv \pi - \partial_0 \phi. \quad (45)$$

Then we have

$$\pi \partial_0 \phi - \mathcal{H} = -\frac{1}{2} \tilde{\pi}^2 + \frac{1}{2} [(\partial_0 \phi)^2 - (\nabla \phi)^2 - m^2 \phi^2] = -\frac{1}{2} \tilde{\pi}^2 + \mathcal{L}. \quad (46)$$

We can thus separate the integration over  $\phi$  from the integration over  $\tilde{\pi}$ : inserting Eq. (46) as well as  $\mu = 0$  into Eq. (41) yields

$$\begin{aligned} Z &= \int \mathcal{D}\tilde{\pi} \exp\left(-\frac{1}{2} \int_X \tilde{\pi}^2(\tau, \mathbf{x})\right) \int \mathcal{D}\phi \exp \int_X \mathcal{L} \\ &= N \int \mathcal{D}\phi \exp \int_X \mathcal{L}, \end{aligned} \quad (47)$$

where we have absorbed the result of the Gaussian momentum integral into an irrelevant constant, and where we have abbreviated

$$\int_X \equiv \int_0^\beta d\tau \int d^3x. \quad (48)$$

[End of 2nd lecture, Oct 13th, 2008.]

It remains to perform the integral over the Lagrangian which can be done exactly for a non-interacting field. We denote four-momenta by capital letters,

$$X \equiv (t, \mathbf{x}) = (-i\tau, \mathbf{x}), \quad K \equiv (k_0, \mathbf{k}) = (-i\omega_n, \mathbf{k}), \quad (49)$$

where  $\omega_n$  are the ‘‘Matsubara frequencies’’ which we explain now. The Fourier transform of the field is

$$\phi(X) = \frac{1}{\sqrt{TV}} \sum_K e^{-iK \cdot X} \phi(K) = \frac{1}{\sqrt{TV}} \sum_K e^{i(\omega_n \tau + \mathbf{k} \cdot \mathbf{x})} \phi(K), \quad (50)$$

with the Minkowski scalar product  $K \cdot X = k_0 x_0 - \mathbf{k} \cdot \mathbf{x}$ . The normalization is chosen such that the Fourier-transformed fields  $\phi(K)$  are dimensionless. We know from the previous section that the field has to be periodic,  $\phi(0, \mathbf{x}) = \phi(\beta, \mathbf{x})$ . To fulfill this periodicity requirement we need  $e^{i\omega_n \beta} = 1$ , i.e.,  $\omega_n \beta$  has to be an integer multiple of  $2\pi$ , or

$$\omega_n = 2\pi nT, \quad n \in \mathbb{Z}. \quad (51)$$

With the Fourier transform (50), and the relation

$$\int_X e^{iK \cdot X} = \frac{V}{T} \delta_{K,0}, \quad (52)$$

we have

$$\begin{aligned} \int_X \mathcal{L} &= -\frac{1}{2} \int_X [(\partial_\tau \phi)^2 + (\nabla \phi)^2 + m^2 \phi^2] \\ &= -\frac{1}{2} \sum_K \phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K), \end{aligned} \quad (53)$$



with the free (hence the subscript “0”) inverse propagator in momentum space

$$D_0^{-1}(K) = \omega_n^2 + k^2 + m^2. \quad (54)$$

Explicitly, we have for example for the first term,

$$\begin{aligned} \int_X (\partial_\tau \phi)^2 &= \frac{1}{TV} \int_X \sum_{K,Q} [\partial_\tau e^{i(\omega_n \tau + \mathbf{k} \cdot \mathbf{x})} \phi(K)] [\partial_\tau e^{i(\omega_m \tau + \mathbf{q} \cdot \mathbf{x})} \phi(Q)] \\ &= -\frac{1}{TV} \int_X \sum_{K,Q} \omega_n \omega_m e^{-i(K+Q) \cdot X} \phi(K) \phi(Q) \\ &= \frac{1}{T^2} \sum_K \omega_n^2 \phi(-K) \phi(K). \end{aligned} \quad (55)$$

Since  $\phi(X)$  is real we have  $\phi(K) = \phi^*(-K)$  and thus

$$Z = N \int \mathcal{D}\phi \exp \left[ -\frac{1}{2} \sum_K \phi^*(K) \frac{D_0^{-1}(K)}{T^2} \phi(K) \right]. \quad (56)$$

We can evaluate this integral by using the general formula

$$\int d^D x e^{-\frac{1}{2} \mathbf{x} \cdot \hat{A} \mathbf{x}} = (2\pi)^{D/2} (\det \hat{A})^{-1/2}, \quad (57)$$

for a hermitian, positive definite matrix  $\hat{A}$ . This identity is a generalization of the one-dimensional gaussian integral

$$\int_{-\infty}^{\infty} e^{-\frac{1}{2} \alpha x^2} = \sqrt{\frac{2\pi}{\alpha}}, \quad (58)$$

and can easily be shown by writing the bilinear  $\mathbf{x} \cdot \hat{A} \mathbf{x}$  in terms of the eigenvalues of  $\hat{A}$  and then using Eq. (58). Consequently,

$$Z = N' \left( \det \frac{D_0^{-1}(K)}{T^2} \right)^{-1/2}, \quad (59)$$

where we have absorbed the constant factor into the new constant  $N'$ , and where the determinant is taken over momentum space (in which the inverse propagator is diagonal). Hence the log of the partition function is, up to a constant,

$$\begin{aligned} \ln Z &= -\frac{1}{2} \ln \det \frac{D_0^{-1}(K)}{T^2} \left( = -\frac{1}{2} \text{Tr} \ln \frac{D_0^{-1}(K)}{T^2} \right) \\ &= -\frac{1}{2} \ln \prod_K \frac{D_0^{-1}(K)}{T^2} \\ &= -\frac{1}{2} \sum_K \ln \frac{D_0^{-1}(K)}{T^2}. \end{aligned} \quad (60)$$

Next we perform the summation over Matsubara frequencies (recall that the sum over  $K$  is a sum over  $k_0 = -i\omega_n$  and over  $\mathbf{k}$ ; the latter will become an integral in the thermodynamic limit).

### A. Summation over bosonic Matsubara frequencies

Here we prove the identity

$$\sum_n \ln \frac{\omega_n^2 + \epsilon_k^2}{T^2} = \frac{\epsilon_k}{T} + 2 \ln \left( 1 - e^{-\epsilon_k/T} \right) + \text{const}, \quad (61)$$

where, in our case,  $\epsilon_k^2 = k^2 + m^2$  (however, for the following calculation we only need that  $\epsilon_k$  is a real number), and where const is an (infinite) number independent of temperature. First, in order to get rid of the log, we write

$$\sum_n \ln \frac{\omega_n^2 + \epsilon_k^2}{T^2} = \int_1^{(\epsilon_k/T)^2} dx^2 \sum_n \frac{1}{(2n\pi)^2 + x^2} + \sum_n \ln[1 + (2n\pi)^2]. \quad (62)$$

We now perform the sum in the integrand which, denoting  $\epsilon_k \equiv Tx$ , we write as a contour integratal,

$$\frac{1}{T} \sum_n \frac{1}{(2n\pi)^2 + x^2} = T \sum_n \frac{1}{\omega_n^2 + \epsilon_k^2} = -\frac{1}{2\pi i} \oint_C d\omega \frac{1}{\omega^2 - \epsilon_k^2} \frac{1}{2} \coth \frac{\omega}{2T}. \quad (63)$$

The second identity follows from the residue theorem,

$$\frac{1}{2\pi i} \oint_C dz f(z) = \sum_n \text{Res } f(z)|_{z=z_n}, \quad (64)$$

where  $z_n$  are the poles of  $f(z)$  in the area enclosed by the contour  $C$ . If we can write the function  $f$  as  $f(z) = \varphi(z)/\psi(z)$ , with analytic functions  $\varphi(z)$ ,  $\psi(z)$ , the residues are

$$\text{Res } f(z)|_{z=z_n} = \frac{\varphi(z_n)}{\psi'(z_n)}. \quad (65)$$

The contour  $C$  in Eq. (63) encloses all poles of  $\coth[\omega/(2T)]$  (and none of  $1/(\omega^2 - \epsilon_k^2)$ ). The denominator of  $\coth[\omega/(2T)]$  is  $e^{\omega/2T} - e^{-\omega/2T}$  which vanishes when  $\omega/2T$  is an integer multiple of  $i\pi$ , i.e., when  $\omega = i\omega_n$  with the Matsubara frequencies  $\omega_n$ . Hence, in the above notation,

$$\begin{aligned} \varphi(\omega) &= \frac{1}{2} \frac{e^{\omega/(2T)} + e^{-\omega/(2T)}}{\omega^2 - \epsilon_k^2}, & \psi(\omega) &= e^{\omega/(2T)} - e^{-\omega/(2T)}, \\ \Rightarrow \frac{\varphi(i\omega_n)}{\psi'(i\omega_n)} &= -T \frac{1}{\omega_n^2 + \epsilon_k^2}, \end{aligned} \quad (66)$$

from which Eq. (63) follows immediately. Next, we deform the contour (which consists of infinitely many circles surrounding the poles) and obtain

$$\begin{aligned} T \sum_n \frac{1}{\omega_n^2 + \epsilon_k^2} &= -\frac{1}{2\pi i} \int_{-i\infty+\eta}^{i\infty+\eta} d\omega \frac{1}{\omega^2 - \epsilon_k^2} \frac{1}{2} \coth \frac{\omega}{2T} - \frac{1}{2\pi i} \int_{i\infty-\eta}^{-i\infty-\eta} d\omega \frac{1}{\omega^2 - \epsilon_k^2} \frac{1}{2} \coth \frac{\omega}{2T} \\ &= -\frac{1}{2\pi i} \int_{-i\infty+\eta}^{i\infty+\eta} d\omega \frac{1}{\omega^2 - \epsilon_k^2} \coth \frac{\omega}{2T}, \end{aligned} \quad (67)$$

where we have changed the integration variable  $\omega \rightarrow -\omega$  in the second integral of the first line. We now use the residue theorem a second time: we can close the contour in the positive half plane and pick up the poles  $\omega = \pm\epsilon_k$  (in our simple case  $\epsilon_k > 0$ , but we can keep the result general in order to use it later for the case of a nonvanishing chemical potential),

$$\begin{aligned} T \sum_n \frac{1}{\omega_n^2 + \epsilon_k^2} &= \Theta(\epsilon_k) \frac{1}{2\epsilon_k} \coth \frac{\epsilon_k}{2T} - \Theta(-\epsilon_k) \frac{1}{2\epsilon_k} \coth \frac{-\epsilon_k}{2T} \\ &= \frac{1}{2\epsilon_k} \coth \frac{\epsilon_k}{2T} \\ &= \frac{1}{2\epsilon_k} [1 + 2f_B(\epsilon_k)], \end{aligned} \quad (68)$$

(note minus sign from clockwise contour integration) with the Bose distribution function

$$f_B(\epsilon) \equiv \frac{1}{e^{\epsilon/T} - 1}. \quad (69)$$

We thus have found

$$\frac{1}{T} \sum_n \frac{1}{(2n\pi)^2 + x^2} = \frac{1}{Tx} \left( \frac{1}{2} + \frac{1}{e^x - 1} \right). \quad (70)$$

Now we insert the result into the original expression (62) and integrate over  $x^2$  to obtain (with const denoting  $T$ -independent constants)

$$\begin{aligned} \sum_n \ln \frac{\omega_n^2 + \epsilon_k^2}{T^2} &= \int_1^{(\epsilon_k/T)^2} dx^2 \frac{1}{x} \left( \frac{1}{2} + \frac{1}{e^x - 1} \right) + \text{const} \\ &= \frac{\epsilon_k}{T} + 2 \ln \left( 1 - e^{-\epsilon_k/T} \right) + \text{const}, \end{aligned} \quad (71)$$

which is the result we wanted to prove.

**Exercise 1:** Show via contour integration that

$$T \sum_{k_0} \frac{1}{[(p_0 - k_0)^2 - \omega_q^2](k_0^2 - \omega_k^2)} = - \sum_{e_1, e_2 = \pm} \frac{e_1 e_2}{4\omega_k \omega_q} \frac{1}{p_0 - e_1 \omega_k - e_2 \omega_q} [1 + f_B(e_1 \omega_k) + f_B(e_2 \omega_q)], \quad (72)$$

with  $k_0 = -i\omega_n$ ,  $p_0 = -i\omega_m$  bosonic Matsubara frequencies and  $\omega_k, \omega_q > 0$ .

## B. Bosonic partition function

Inserting the result from the Matsubara sum into Eq. (60) and taking the thermodynamic limit yields the (log of) the bosonic partition function,

$$\ln Z = -V \int \frac{d^3 k}{(2\pi)^3} \left[ \frac{\epsilon_k}{2T} + \ln \left( 1 - e^{-\epsilon_k/T} \right) \right]. \quad (73)$$

## V. COMPLEX NON-INTERACTING SCALAR FIELD

Next we discuss a complex bosonic field. Although we still neglect interactions, this will already lead to new physics compared to the real field, namely Bose-Einstein condensation. We start from the Lagrangian

$$\mathcal{L}_0 = \partial_\mu \phi^* \partial^\mu \phi - m^2 |\phi|^2 - \lambda |\phi|^4. \quad (74)$$

We set the coupling to zero,  $\lambda = 0$ . We added the subscript “0” to indicate that the chemical potential is not included yet. In contrast to the previous case, we now have a conserved charge and thus can introduce a chemical potential. To this end, we first identify the conserved current. We know from Noether’s theorem that the conserved current is related to the symmetry of the Lagrangian. We see that  $\mathcal{L}$  is invariant under  $U(1)$  rotations of the field,

$$\phi \rightarrow e^{-i\alpha} \phi. \quad (75)$$

To find the current we formally extend this symmetry to a local symmetry  $\alpha(x)$  and transform the Lagrangian,

$$\mathcal{L}_0 \rightarrow \mathcal{L}_0 + |\phi|^2 \partial_\mu \alpha \partial^\mu \alpha + i \partial_\mu \alpha (\phi^* \partial^\mu \phi - \phi \partial^\mu \phi^*). \quad (76)$$

Now we write down the equation of motion for  $\alpha$ . We see that the transformed Lagrangian does not depend on  $\alpha$ , but only on its derivative. Consequently, the quantity

$$\frac{\partial \mathcal{L}_0}{\partial (\partial_\mu \alpha)} = |\phi|^2 \partial_\mu \alpha + i(\phi^* \partial_\mu \phi - \phi \partial_\mu \phi^*) \quad (77)$$

is conserved. If we now go back to constant  $\alpha$  we see that we have the conserved current

$$j_\mu \equiv i(\phi^* \partial_\mu \phi - \phi \partial_\mu \phi^*), \quad \partial^\mu j_\mu = 0. \quad (78)$$

The conserved charge (density) is thus

$$j_0 = i(\phi^* \partial_0 \phi - \phi \partial_0 \phi^*). \quad (79)$$

This is needed to introduce a chemical potential  $\mu$ . In the following we want to see how the chemical potential enters the Lagrangian. We will see that we cannot simply add a term  $\mu j_0$  to  $\mathcal{L}_0$ . We rather have to look at the expression

$\pi\partial_0\phi - \mathcal{H} + \mu\mathcal{N}$  in the action. To this end, we first introduce the two real fields  $\phi_1, \phi_2$  as real and imaginary parts of  $\phi$ ,

$$\phi = \frac{1}{\sqrt{2}}(\phi_1 + i\phi_2). \quad (80)$$

Then, the Lagrangian becomes

$$\mathcal{L}_0 = \frac{1}{2} [\partial_\mu\phi_1\partial^\mu\phi_1 + \partial_\mu\phi_2\partial^\mu\phi_2 - m^2(\phi_1^2 + \phi_2^2)]. \quad (81)$$

The conjugate momenta are

$$\pi_i = \frac{\partial\mathcal{L}_0}{\partial(\partial_0\phi_i)} = \partial_0\phi_i, \quad i = 1, 2. \quad (82)$$

Consequently, with  $j_0 = \phi_2\pi_1 - \phi_1\pi_2$  which follows from Eqs. (79), (80), and (82), we have

$$\begin{aligned} \mathcal{H} - \mu\mathcal{N} &= \pi_1\partial_0\phi_1 + \pi_2\partial_0\phi_2 - \mathcal{L}_0 - \mu\mathcal{N} \\ &= \frac{1}{2} [\pi_1^2 + \pi_2^2 + (\nabla\phi_1)^2 + (\nabla\phi_2)^2 + m^2(\phi_1^2 + \phi_2^2)] - \mu(\phi_2\pi_1 - \phi_1\pi_2). \end{aligned} \quad (83)$$

For the partition function, see Eq. (41), we again introduce a shifted momentum field, as we did in the previous section. This time we have to include the chemical potential in the shift in order to separate the field integration from the momentum integration,

$$\tilde{\pi}_1 \equiv \pi_1 - \partial_0\phi_1 - \mu\phi_2, \quad \tilde{\pi}_2 \equiv \pi_2 - \partial_0\phi_2 + \mu\phi_1. \quad (84)$$

This yields

$$\pi_1\partial_0\phi_1 + \pi_2\partial_0\phi_2 - \mathcal{H} + \mu\mathcal{N} = -\frac{1}{2}(\tilde{\pi}_1^2 + \tilde{\pi}_2^2) + \mathcal{L}, \quad (85)$$

where the new Lagrangian  $\mathcal{L}$  now includes the chemical potential,

$$\mathcal{L} = \frac{1}{2} [(\partial_0\phi_1)^2 + (\partial_0\phi_2)^2 - (\nabla\phi_1)^2 - (\nabla\phi_2)^2 + (\mu^2 - m^2)(\phi_1^2 + \phi_2^2) + 2\mu(\phi_2\partial_0\phi_1 - \phi_1\partial_0\phi_2)]. \quad (86)$$

In terms of the complex field  $\phi$ , the Lagrangian reads

$$\mathcal{L} = |(\partial_0 - i\mu)\phi|^2 - |\nabla\phi|^2 - m^2|\phi|^2. \quad (87)$$

Thus we see that the effect of the chemical potential is to add, besides the expected term  $\mu j_0$ , the additional term  $\mu^2(\phi_1^2 + \phi_2^2)/2$ .

[End of 3rd lecture, Oct 20th, 2008.]

In order to compute the partition function, we Fourier transform the fields  $\phi_1, \phi_2$  as discussed for the scalar field. However, anticipating Bose-Einstein condensation, we separate the zero-momentum mode  $\zeta_i = \phi_i(K=0)$ ,

$$\phi_i(X) = \zeta_i + \frac{1}{\sqrt{TV}} \sum_{K \neq 0} e^{-iK \cdot X} \phi(K). \quad (88)$$

The condensate  $\zeta_i$  plays the role of a vacuum expectation value of the field. It breaks the  $U(1)$  symmetry spontaneously. We can choose any of the degenerate directions in the complex plane, for instance  $\zeta_2 = 0$  and will denote  $\zeta \equiv \zeta_1$ . With the Lagrangian (86) the action then becomes

$$\int_X \mathcal{L} = \frac{V}{T} \frac{\mu^2 - m^2}{2} \zeta^2 - \frac{1}{2} \sum_K (\phi_1(-K), \phi_2(-K)) \frac{D_0^{-1}}{T^2} \begin{pmatrix} \phi_1(K) \\ \phi_2(K) \end{pmatrix}, \quad (89)$$

with the  $2 \times 2$  inverse propagator

$$D_0^{-1}(K) = \begin{pmatrix} -K^2 + m^2 - \mu^2 & -2i\mu k_0 \\ 2i\mu k_0 & -K^2 + m^2 - \mu^2 \end{pmatrix}. \quad (90)$$

In deriving the action (89) we have used that the integrals over mixed terms, i.e., over a product of the condensate  $\zeta$  and the momentum sum (excluding the mode  $K = 0$ ), vanish. We see that the chemical potential induces off-diagonal terms in the propagator. Now from the partition function

$$Z = N \int \mathcal{D}\phi_1 \mathcal{D}\phi_2 \exp \int_X \mathcal{L}, \quad (91)$$

we obtain, dropping the constant terms,

$$\begin{aligned} \ln Z &= \frac{V}{T} \frac{\mu^2 - m^2}{2} \zeta^2 - \frac{1}{2} \ln \left( \det \frac{D_0^{-1}(K)}{T^2} \right) \\ &= \frac{V}{T} \frac{\mu^2 - m^2}{2} \zeta^2 - \frac{1}{2} \ln \prod_K \frac{1}{T^4} [(-K^2 + m^2 - \mu^2)^2 - 4\mu^2 k_0^2] \\ &= \frac{V}{T} \frac{\mu^2 - m^2}{2} \zeta^2 - \frac{1}{2} \ln \prod_K \frac{1}{T^4} [(\epsilon_k - \mu)^2 - k_0^2][(\epsilon_k + \mu)^2 - k_0^2] \\ &= \frac{V}{T} \frac{\mu^2 - m^2}{2} \zeta^2 - \frac{1}{2} \sum_K \left[ \ln \frac{(\epsilon_k - \mu)^2 - k_0^2}{T^2} + \ln \frac{(\epsilon_k + \mu)^2 - k_0^2}{T^2} \right], \end{aligned} \quad (92)$$

where we defined

$$\epsilon_k \equiv \sqrt{k^2 + m^2}. \quad (93)$$

We can now use the result of the Matsubara summation from above, Eq. (61), to obtain

$$\ln Z = \frac{V}{T} \frac{\mu^2 - m^2}{2} \zeta^2 - V \int \frac{d^3 k}{(2\pi)^3} \left[ \frac{\epsilon_k}{T} + \ln \left( 1 - e^{-(\epsilon_k - \mu)/T} \right) + \ln \left( 1 - e^{-(\epsilon_k + \mu)/T} \right) \right]. \quad (94)$$

This gives the thermodynamic potential

$$\frac{\Omega}{V} = \frac{m^2 - \mu^2}{2} \zeta^2 + T \int \frac{d^3 k}{(2\pi)^3} \left[ \ln \left( 1 - e^{-(\epsilon_k - \mu)/T} \right) + \ln \left( 1 - e^{-(\epsilon_k + \mu)/T} \right) \right], \quad (95)$$

where we have dropped the (infinite) vacuum contribution, i.e., we actually consider  $\Omega - \Omega_{T=0}$ . (Note that  $\lim_{T \rightarrow 0} T \ln(1 - e^{-E/T}) = -E\Theta(-E)$ .) The condensate  $\zeta$  has to be determined from minimizing the potential,

$$0 = \frac{\partial \Omega}{\partial \zeta} = (m^2 - \mu^2) \zeta. \quad (96)$$

We see that  $\zeta = 0$  for  $|\mu| < m$ . In this case, there is no Bose condensation and all particles sit in the thermal states. For  $|\mu| = m$ ,  $\zeta$  remains undetermined. This is due to our neglecting the interactions. From usual  $\phi^4$  theory at zero temperature we know that the interactions may lead to a nonvanishing vacuum expectation value (“mexican hat potential”). But for now we have dropped the  $\phi^4$  term for simplicity. In this case, we can determine  $\zeta$  by fixing the density. This may or may not correspond to the physical situation we are interested in. In many nonrelativistic applications indeed the density is fixed. However, for example in quark matter in neutron stars the chemical potential typically is fixed, not the charge. In this case we need to include interactions to fix the condensate (of course the noninteracting limit may be a bad approximation for other reasons too). The charge density is

$$Q = -\frac{\partial \Omega}{\partial \mu} = \mu \zeta^2 + \sum_{e=\pm} e \int \frac{d^3 k}{(2\pi)^3} \frac{1}{e^{(\epsilon_k - e\mu)/T} - 1}. \quad (97)$$

Before discussing the role of the condensate  $\zeta$  let us approximate the thermal part for small and large temperatures. We first introduce the new integration variable  $x = k/T$  to obtain

$$Q = \frac{T^3}{2\pi^2} \int_0^\infty dx x^2 \left( \frac{1}{e^{\sqrt{x^2 + (m/T)^2 - \mu/T} - 1} - 1} - \frac{1}{e^{\sqrt{x^2 + (m/T)^2 + \mu/T} - 1} - 1} \right). \quad (98)$$

Now we expand for small  $T$ ,  $\sqrt{x^2 + (m/T)^2} \simeq m/T + Tx^2/(2m)$ . Then, with the new integration variable  $y = \sqrt{T/(2m)}x$ , we have

$$\begin{aligned} Q &\simeq \frac{T^3}{2\pi^2} \int_0^\infty dx x^2 \left( \frac{e^{-(m-\mu)/T}}{e^{Tx^2/(2m)}} - \frac{e^{-(m+\mu)/T}}{e^{Tx^2/(2m)}} \right) \\ &= \left( \frac{2m}{T} \right)^{3/2} \frac{T^3}{2\pi^2} \left( e^{-(m-\mu)/T} - e^{-(m+\mu)/T} \right) \int_0^\infty dy y^2 e^{-y^2} \\ &= \frac{m^{3/2} T^{3/2}}{2\sqrt{2}\pi^{3/2}} \left( e^{-(m-\mu)/T} - e^{-(m+\mu)/T} \right), \end{aligned} \quad (99)$$

where we have used that the remaining  $y$ -integral evaluates to  $\sqrt{\pi}/4$ . We see that the density is exponentially suppressed for small temperatures. This exponential suppression for massive particles is also typical for other quantities such as the specific heat.

For large temperatures we can use Eq. (98) and neglect the terms  $m/T$  and  $\mu/T$  in the integrand to obtain

$$Q_+ = Q_- = \frac{T^3}{2\pi^2} \int_0^\infty dx \frac{x^2}{e^x - 1} = \frac{\zeta(3)T^3}{\pi^2}, \quad (100)$$

where  $Q_+$  and  $Q_-$  are the particle and antiparticle contributions, respectively. We see that they become identical for large  $T$  and thus the total charge  $Q = Q_+ - Q_-$  vanishes. This is easy to understand: the difference in energies between particles and antiparticles is  $2\mu$  for all momenta. If  $T$  is sufficiently large, i.e.,  $T \gg \mu$ , then this difference is not “resolved” and particle and antiparticle states become practically equally populated. For  $T$  of the order of  $\mu$  or smaller, the chemical potential induces an asymmetry between particles and antiparticles, favoring particles for  $\mu > 0$  and antiparticles for  $\mu < 0$ .

**Exercise 2:** Compute the specific heat (at constant chemical potential)  $c_V = T\partial S/\partial T$ , where  $S = -\partial\Omega/\partial T$  is the entropy, and find analytic approximations for the limits of small and large temperatures.

Let us now discuss the condensate. There is a zero-temperature contribution  $\mu\zeta^2$  coming from the bosons in the zero-momentum state. For a given density, the system populates as many thermal states as possible until there is no more “space”. Note that the contribution of the thermal integral is bounded with its maximum at  $\mu^2 = m^2$ . This maximum value defines a critical density for a given temperature  $T$ . For densities larger than this critical density, the condensate gets populated. The population is “macroscopic”, i.e., proportional to the volume. The value of the condensate is given by

$$\zeta^2 = \frac{1}{m} \left( Q - \sum_{\epsilon=\pm} e \int \frac{d^3k}{(2\pi)^3} \frac{1}{e^{(\epsilon_k - \epsilon m)/T} - 1} \right). \quad (101)$$

The critical temperature  $T_c$  for a given charge density  $Q$  is then given by the implicit equation

$$Q = \sum_{\epsilon=\pm} e \int \frac{d^3k}{(2\pi)^3} \frac{1}{e^{(\epsilon_k - \epsilon m)/T_c} - 1}. \quad (102)$$

In the nonrelativistic limit we have  $\epsilon_k = k^2/(2m)$  and condensation occurs for  $\mu = 0$ . In this case,  $T_c$  can be computed as

$$\begin{aligned} Q &= \int \frac{d^3k}{(2\pi)^3} \frac{1}{e^{k^2/(2mT_c)} - 1} \\ &= \frac{1}{2\pi^2} \int_0^\infty dk k^2 \frac{1}{e^{k^2/(2mT_c)} - 1} \\ &= \frac{(2mT_c)^{3/2}}{2\pi^2} \int_0^\infty dx \frac{x^2}{e^{x^2} - 1} \\ &= \frac{(2mT_c)^{3/2}}{2\pi^2} \frac{\sqrt{\pi}}{4} \zeta(3/2), \end{aligned} \quad (103)$$

which implies

$$T_c = \frac{2\pi}{m} \left( \frac{Q}{\zeta(3/2)} \right)^{2/3}. \quad (104)$$

In the ultrarelativistic limit it is instructive to compute particle and antiparticle contributions separately. With  $\mu = m$  and  $\epsilon_k \pm m \simeq k \pm m$  we have

$$Q_{\pm} \simeq \frac{T_c^3}{2\pi^2} \int_0^{\infty} dx \frac{x^2}{e^{x \mp m/T_c} - 1}. \quad (105)$$

Up to first order in  $m/T_c$  we have

$$\frac{1}{e^{x \mp m/T_c} - 1} = \frac{e^{\pm m/T_c}}{e^x - e^{\pm m/T_c}} \simeq \frac{1}{e^x - 1} \left( 1 \pm \frac{m}{T_c} \right) \pm \frac{m/T_c}{(e^x - 1)^2}. \quad (106)$$

Consequently,

$$\begin{aligned} Q_{\pm} &\simeq \frac{T_c^3}{2\pi^2} \int_0^{\infty} dx \frac{x^2}{e^x - 1} \pm \frac{T_c^2 m}{2\pi^2} \left[ \int_0^{\infty} dx \frac{x^2}{e^x - 1} + \int_0^{\infty} dx \frac{x^2}{(e^x - 1)^2} \right] \\ &= \frac{T_c^3}{2\pi^2} \left[ 2\zeta(3) \pm \frac{m\pi^2}{3T_c} \right]. \end{aligned} \quad (107)$$

We see that the antiparticles have an interesting effect: did we neglect  $Q_-$ , the critical temperature would be  $T_c \propto Q_+^{1/3}$ . In the sum of both contributions, however, the leading term cancels and we get the very different result

$$T_c = \sqrt{\frac{3Q}{m}}. \quad (108)$$

Bose-Einstein condensation is a phenomenon occurring in a huge variety of systems. It was first observed with bosonic atoms in 1995, awarded with the Nobel prize 2001. It often has spectacular phenomenological consequences, such as in superfluid He-4. It can also occur for excitons in semiconductors, and for mesons such as pions and kaons in neutron stars. One can even think of superconductivity in fermionic systems as a Bose-Einstein condensate, since Cooper pairs of fermions can be viewed as bosons. Very recent experiments have shown that this picture indeed is valid, i.e., there is a crossover from a superfluid (at weak coupling) to a Bose-Einstein condensate (at strong coupling), not a phase transition.

## VI. NON-INTERACTING FERMIONS

We shall now turn to fermions and compute their partition function. We shall see that there are two important differences to the bosonic case. Firstly, the fields over which we integrate in the functional integral are anticommuting, which yields a different result for the functional integration. Secondly, we shall have antiperiodicity instead of periodicity in the fields, which yields different Matsubara frequencies. Both differences are related to the Pauli principle.

### A. Grassmann Algebra and antiperiodicity in $\beta$ for fermion fields

We start by defining the so-called Grassmann Algebra: on an  $r$ -dimensional vector space with basis vectors  $\eta_1, \dots, \eta_r$  we define an anticommuting product

$$\eta_i \eta_j = -\eta_j \eta_i, \quad (109)$$

to obtain the Grassmann Algebra  $\mathcal{A}$ . The algebra has  $2^r$  basis elements  $1, \eta_i, \eta_i \eta_j, \dots, \eta_1 \eta_2 \dots \eta_r$ . Note that Eq. (109) implies  $\eta_i^2 = 0$ . One needs a sign convention to define the derivatives on this space. For example, for  $j \neq k$ ,

$$\frac{\partial}{\partial \eta_j} \eta_j \eta_k = \eta_k, \quad \frac{\partial}{\partial \eta_k} \eta_j \eta_k = -\eta_j. \quad (110)$$

[End of 4th lecture, Oct 27th, 2008]

This is a convenient convention since one can think of the derivative operator as anticommuting with the variable itself. Second derivatives of any product of  $\eta$ 's vanish (they vanish if there is at most one factor of the variable with respect to which the derivative is taken; if there are two factors the product itself vanishes). This already shows

that integration on the Grassmann space is a bit different than one is used to: since the differential operator squared vanishes, the integration cannot be the inverse operation to differentiation. We require the integral to be linear and translationally invariant. In a one-dimensional space this means

$$\int d\eta f(\eta) = \int d\eta f(\eta + \zeta), \quad \int d\eta (a\eta + b) = a \int d\eta \eta + b \int d\eta, \quad (111)$$

with  $a, b$  complex numbers and  $\zeta \in \mathcal{A}$ . Here, the integration range is always the whole space, i.e., we can only talk about “definite” integrals; “indefinite” integrals, as one is used to from  $c$ -numbers do not exist (since this would be an operation inverse to differentiation). If we define functions via their Taylor expansion,  $f$  must have the form  $f(\eta) = a\eta + b$ . Then, from (111) we conclude

$$\int d\eta = 0, \quad \int d\eta \eta = 1, \quad (112)$$

where the second is a normalization (not a consequence from (111)). Equipped with these properties we can turn to fermions.

First we consider a simple system with two states  $|0\rangle$  and  $|1\rangle$ , and creation and annihilation operators  $a$  and  $a^\dagger$  which obey the anticommutation relations

$$\{a, a^\dagger\} = 1, \quad a^2 = (a^\dagger)^2 = 0. \quad (113)$$

Moreover we consider the Grassmann Algebra generated by the two variables  $\eta$  and  $\eta^*$  (these shall correspond to the fermion fields later), and the states

$$|\eta\rangle \equiv e^{-\eta a^\dagger} |0\rangle = (1 - \eta a^\dagger) |0\rangle = |0\rangle - \eta |1\rangle, \quad (114a)$$

$$\langle \eta | \equiv \langle 0 | e^{-a \eta^*} = \langle 0 | (1 - a \eta^*) = \langle 0 | - \langle 1 | \eta^*. \quad (114b)$$

We assume that  $a$  and  $\eta$  anticommute such that  $a|\eta\rangle = \eta|0\rangle$  and thus  $a|\eta\rangle = \eta|\eta\rangle$ . We shall need this relation later. We also need

$$\langle \eta | 0 \rangle = \langle 0 | \eta \rangle = 1, \quad \langle 1 | \eta \rangle = \langle \eta | 1 \rangle^* = -\eta, \quad (115)$$

which is obvious from Eqs. (114), and

$$\langle \eta_1 | \eta_2 \rangle = e^{\eta_1^* \eta_2}, \quad (116)$$

which follows from inserting  $1 = |0\rangle\langle 0| + |1\rangle\langle 1|$  and using Eqs. (115). Also, with Eqs. (114) and the rules for integration (112) (generalized to two dimensions) we find

$$\begin{aligned} \int d\eta^* d\eta e^{-\eta^* \eta} |\eta\rangle\langle \eta| &= \int d\eta^* d\eta (1 - \eta^* \eta) (|0\rangle\langle 0| - \eta |1\rangle\langle 0| - |0\rangle\langle 1| \eta^* + |1\rangle\langle 1| \eta \eta^*) \\ &= |0\rangle\langle 0| + |1\rangle\langle 1| = 1. \end{aligned} \quad (117)$$

And, finally, upon inserting unity twice and using Eqs. (115)

$$\begin{aligned} \int d\eta^* d\eta e^{-\eta^* \eta} \langle -\eta | A | \eta \rangle &= \int d\eta^* d\eta (1 - \eta^* \eta) (\langle 0 | A | 0 \rangle + \eta^* \langle 1 | A | 0 \rangle - \eta \langle 0 | A | 1 \rangle - \eta^* \eta \langle 1 | A | 1 \rangle) \\ &= \langle 0 | A | 0 \rangle + \langle 1 | A | 1 \rangle = \text{Tr } A. \end{aligned} \quad (118)$$

From this formula we compute the partition function for the Hamiltonian  $\hat{H} = \omega a^\dagger a$ ,

$$Z = \text{Tr} e^{-\beta \hat{H}} = \int d\eta^* d\eta e^{-\eta^* \eta} \langle -\eta | e^{-\beta \hat{H}} | \eta \rangle. \quad (119)$$

The important difference to the bosonic case can already be seen here, namely the  $-\eta$  as the final state of the transition amplitude. We can now proceed analogously to the bosonic case by dividing the “time” interval into  $N$  pieces of width  $\Delta t$  and inserting unity from Eq. (117)  $N - 1$  times. We obtain

$$Z = \int_{\eta(\beta) = -\eta(0)} \mathcal{D}\eta^* \mathcal{D}\eta \exp \left( - \int_0^\beta d\tau [\eta^* \partial_\tau \eta + H(\eta^*, \eta)] \right). \quad (120)$$

Before we generalize this to the case of Dirac fields let us discuss the fermionic Lagrangian.

[\[End of 5th lecture, Nov 3rd, 2008\]](#)



### B. Fermionic Lagrangian and conserved charge

We start with the non-interacting Lagrangian

$$\mathcal{L}_0 = \bar{\psi} (i\gamma^\mu \partial_\mu - m) \psi, \quad (121)$$

where  $\bar{\psi} = \psi^\dagger \gamma^0$ , and where the Dirac matrices are given in the Dirac representation by

$$\gamma^0 = \begin{pmatrix} 1 & 0 \\ 0 & -1 \end{pmatrix}, \quad \gamma^i = \begin{pmatrix} 0 & \sigma_i \\ -\sigma_i & 0 \end{pmatrix}, \quad (122)$$

with the Pauli matrices  $\sigma_i$ . The general properties of the Dirac matrices are

$$\{\gamma^\mu, \gamma^\nu\} = 2g^{\mu\nu}, \quad (\gamma^0)^2 = 1, \quad (\gamma^i)^2 = -1, \quad (\gamma^0)^\dagger = \gamma^0, \quad (\gamma^i)^\dagger = -\gamma^i, \quad (123)$$

where  $g^{\mu\nu}$  is the Minkowski metric.

As for the bosons we are interested in the theory with a chemical potential. To this end, we determine the conserved current with the same method as above. The Lagrangian is invariant under the transformation  $\psi \rightarrow e^{-i\alpha} \psi$ . Considering a local transformation  $\alpha(x)$ , we have

$$\mathcal{L}_0 \rightarrow \mathcal{L}_0 + \bar{\psi} \gamma^\mu (\partial_\mu \alpha) \psi. \quad (124)$$

From the equation of motion for  $\alpha$  we then conclude that the current

$$j^\mu = \frac{\partial \mathcal{L}}{\partial (\partial_\mu \alpha)} = \bar{\psi} \gamma^\mu \psi \quad (125)$$

is conserved, i.e.,

$$\partial_\mu j^\mu = 0, \quad (126)$$

and the conserved charge (density) is given by

$$Q = \psi^\dagger \psi. \quad (127)$$

The conjugate momentum is

$$\pi = \frac{\partial \mathcal{L}}{\partial (\partial_0 \psi)} = i\psi^\dagger. \quad (128)$$

We see that we have to treat  $\psi$  and  $\psi^\dagger$  as independent variables, in accordance to what we have discussed before in terms of  $\eta$  and  $\eta^*$ . Consequently, the Hamiltonian becomes

$$\mathcal{H} = \pi \partial_0 \psi - \mathcal{L} = \bar{\psi} (i\boldsymbol{\gamma} \cdot \nabla + m) \psi. \quad (129)$$

Here and in the following we mean by the scalar product  $\boldsymbol{\gamma} \cdot \nabla$  the product where the Dirac matrices appear with a lower index  $\gamma_i$ , i.e., the negative of the  $\gamma^i$  given in Eq. (122).

### C. Partition function for fermions

Now we recall that for the partition function we need  $i\pi \partial_\tau \psi - \mathcal{H} + \mu \mathcal{N}$  (see for instance Eq. (41)). With the Hamiltonian (129) and the generalization of the fermionic partition function (120) to fields  $\psi$ ,  $\psi^\dagger$  we obtain

$$Z = \int_{\text{antiperiodic}} \mathcal{D}\psi^\dagger \mathcal{D}\psi \exp \left[ \int_X \bar{\psi} (-\gamma^0 \partial_\tau - i\boldsymbol{\gamma} \cdot \nabla + \gamma^0 \mu - m) \psi \right]. \quad (130)$$

Analogous to the bosonic case, we introduce the Fourier transform (note different dimensionality of fields compared to bosons; here the field has mass dimension 3/2)

$$\psi(X) = \frac{1}{\sqrt{V}} \sum_K e^{-iK \cdot X} \psi(K), \quad \bar{\psi}(X) = \frac{1}{\sqrt{V}} \sum_K e^{iK \cdot X} \bar{\psi}(K), \quad \int_X e^{iK \cdot X} = \frac{V}{T} \delta_{K,0}, \quad (131)$$

again with  $k_0 = -i\omega_n$  such that  $K \cdot X = -(\omega_n\tau + \mathbf{k} \cdot \mathbf{x})$ . Now antiperiodicity requires  $\psi(0, \mathbf{x}) = -\psi(\beta, \mathbf{x})$ , which implies  $e^{i\omega_n\beta} = -1$  and thus the fermionic Matsubara frequencies are

$$\omega_n = (2n + 1)\pi T. \quad (132)$$

With the Fourier decomposition we find

$$\int_X \bar{\psi} (-\gamma^0 \partial_\tau - i\boldsymbol{\gamma} \cdot \nabla + \gamma^0 \mu - m) \psi = - \sum_K \psi^\dagger(K) \frac{G_0^{-1}(K)}{T} \psi(K), \quad (133)$$

with the free inverse fermion propagator in momentum space

$$G_0^{-1}(K) = -\gamma^\mu K_\mu - \gamma^0 \mu + m. \quad (134)$$

Before we continue let us introduce a useful form of the inverse propagator in terms of energy projectors. This form will not be needed here but is very helpful for more difficult calculations. In particular it allows inversion in a simple way. We can write

$$G_0^{-1}(K) = - \sum_{\epsilon=\pm} (k_0 + \mu - \epsilon\epsilon_k) \gamma^0 \Lambda_{\mathbf{k}}^\epsilon, \quad (135)$$

where the projectors onto positive and negative energy states are given by

$$\Lambda_{\mathbf{k}}^\epsilon \equiv \frac{1}{2} \left( 1 + e\gamma^0 \frac{\boldsymbol{\gamma} \cdot \mathbf{k} + m}{\epsilon_k} \right). \quad (136)$$

These (hermitian) projectors are complete and orthogonal,

$$\Lambda_{\mathbf{k}}^+ + \Lambda_{\mathbf{k}}^- = 1, \quad \Lambda_{\mathbf{k}}^\epsilon \Lambda_{\mathbf{k}}^{\epsilon'} = \delta_{\epsilon, \epsilon'} \Lambda_{\mathbf{k}}^\epsilon. \quad (137)$$

The first property is trivial to see, the second follows with the anticommutation property  $\{\gamma^0, \gamma^i\} = 0$  which follows from the general anticommutation property in Eq. (123) and with  $(\boldsymbol{\gamma} \cdot \mathbf{k})^2 = -1$ .

From the form of the inverse propagator (135) we can immediately read off the propagator itself,

$$G_0(K) = - \sum_{\epsilon=\pm} \frac{\Lambda_{\mathbf{k}}^\epsilon \gamma^0}{k_0 + \mu - \epsilon\epsilon_k}. \quad (138)$$

With the properties (137) one easily checks that  $G_0^{-1}G_0 = 1$ . One can also rewrite (138) as

$$G_0(K) = \frac{-\gamma^\mu K_\mu - \gamma^0 \mu - m}{(k_0 + \mu)^2 - \epsilon_k^2}. \quad (139)$$

Now let us come back to the partition function. For the functional integration we use

$$\int \prod_k^N d\eta_k^\dagger d\eta_k \exp \left( - \sum_{i,j} \eta_i^\dagger D_{ij} \eta_j \right) = \det D. \quad (140)$$

**Exercise 3:** *Prove this relation by using the above properties of the Grassmann variables.*

Note the difference of this Grassmann integration for fermions with the corresponding formula for bosons (57). We obtain for the partition function

$$\begin{aligned} Z &= \int_{\text{antiperiodic}} \mathcal{D}\psi^\dagger \mathcal{D}\psi \exp \left[ - \sum_K \psi^\dagger(K) \frac{G_0^{-1}(K)}{T} \psi(K) \right] \\ &= \det \frac{G_0^{-1}(K)}{T} \\ &= \det \frac{1}{T} \begin{pmatrix} -(k_0 + \mu) + m & -\boldsymbol{\sigma} \cdot \mathbf{k} \\ \boldsymbol{\sigma} \cdot \mathbf{k} & (k_0 + \mu) + m \end{pmatrix}, \end{aligned} \quad (141)$$

where the determinant is taken over Dirac space and momentum space. We can use the general formula

$$\det \begin{pmatrix} A & B \\ C & D \end{pmatrix} = \det(AD - BD^{-1}CD), \quad (142)$$

for matrices  $A, B, C, D$  with  $D$  invertible, to get

$$\det G_0^{-1}(K) = \prod_K \left( \frac{k^2 + m^2 - (k_0 + \mu)^2}{T^2} \right)^2. \quad (143)$$

Here we have used  $(\boldsymbol{\sigma} \cdot \mathbf{k})^2 = k^2$ . Consequently,

$$\ln Z = \sum_K \ln \left( \frac{\epsilon_k^2 - (k_0 + \mu)^2}{T^2} \right)^2, \quad \epsilon_k \equiv \sqrt{k^2 + m^2}. \quad (144)$$

With  $k_0 = -i\omega_n$  we can write this as

$$\begin{aligned} \ln Z &= \sum_K \ln \left( \frac{\epsilon_k^2 + (\omega_n + i\mu)^2}{T^2} \right)^2 \\ &= \sum_K \left( \ln \frac{\epsilon_k^2 + (\omega_n + i\mu)^2}{T^2} + \ln \frac{\epsilon_k^2 + (-\omega_n + i\mu)^2}{T^2} \right) \\ &= \sum_K \left( \ln \frac{\omega_n^2 + (\epsilon_k - \mu)^2}{T^2} + \ln \frac{\omega_n^2 + (\epsilon_k + \mu)^2}{T^2} \right), \end{aligned} \quad (145)$$

where, in the second step, we have replaced  $\omega_n$  by  $-\omega_n$  which does not change the result since we sum over all  $n \in \mathbb{Z}$ . Then, the third step can be easily checked by multiplying out all terms.

#### D. Summation over fermionic Matsubara frequencies

We have written the log of the fermionic partition function in a form which is identical to the bosonic one, compare Eq. (145) with Eq. (92). The only difference is the form of the Matsubara frequencies. We can thus compute the sum over fermionic Matsubara frequencies analogous to the sum over bosonic ones, explained in Sec. IV A. As above, we write

$$\sum_n \ln \frac{\omega_n^2 + \epsilon_k^2}{T^2} = \int_1^{(\epsilon_k/T)^2} dx^2 \sum_n \frac{1}{(2n+1)^2\pi^2 + x^2} + \sum_n \ln[1 + (2n+1)^2\pi^2]. \quad (146)$$

And as above, we write the sum as a contour integral, this time with the tanh instead of the coth,

$$\frac{1}{T} \sum_n \frac{1}{(2n+1)^2\pi^2 + x^2} = T \sum_n \frac{1}{\omega_n^2 + \epsilon_k^2} = -\frac{1}{2\pi i} \oint_C d\omega \frac{1}{\omega^2 - \epsilon_k^2} \frac{1}{2} \tanh \frac{\omega}{2T}. \quad (147)$$

(We have denoted  $\epsilon_k \equiv xT$ .) The contour  $C$  encloses all poles of the tanh (and none of  $\frac{1}{\omega^2 - \epsilon_k^2}$ ). The poles of the tanh are given by the zeros of  $e^{\omega/(2T)} + e^{-\omega/(2T)}$ , i.e.,  $\omega/(2T)$  must be an odd integer multiple of  $i\pi/2$ . Therefore, the poles are located at  $i$  times the fermionic Matsubara frequencies,  $\omega = i\omega_n$ . Then, with the residue theorem and with

$$\left( e^{\omega/(2T)} - e^{-\omega/(2T)} \right) \Big|_{\omega=i\omega_n} = 2i(-1)^n, \quad \frac{d}{d\omega} \left( e^{\omega/(2T)} + e^{-\omega/(2T)} \right) \Big|_{\omega=i\omega_n} = \frac{i(-1)^n}{T}, \quad (148)$$

one sees Eq. (147). We can then proceed as for bosons, i.e., we close the contour in the positive half-plane to obtain with the residue theorem

$$\begin{aligned} T \sum_n \frac{1}{\omega_n^2 + \epsilon_k^2} &= -\frac{1}{2\pi i} \int_{-i\infty+\eta}^{i\infty+\eta} d\omega \frac{1}{\omega^2 - \epsilon_k^2} \frac{1}{2} \tanh \frac{\omega}{2T} \\ &= \frac{1}{2\epsilon_k} \tanh \frac{\epsilon_k}{2T} \\ &= \frac{1}{2\epsilon_k} [1 - 2f_F(\epsilon_k)], \end{aligned} \quad (149)$$

where

$$f_F(\epsilon) \equiv \frac{1}{e^{\epsilon/T} - 1}. \quad (150)$$

is the Fermi distribution function. Inserting this result into the original expression (146) yields

$$\begin{aligned} \sum_n \ln \frac{\omega_n^2 + \epsilon_k^2}{T^2} &= T \int_1^{(\epsilon_k/T)^2} dx^2 \frac{1}{2\epsilon_k} [1 - 2f(\epsilon_k)] + \text{const} \\ &= \int_1^{(\epsilon_k/T)^2} dx^2 \frac{1}{x} \left( \frac{1}{2} - \frac{1}{e^x + 1} \right) + \text{const} \\ &= \frac{\epsilon_k}{T} + 2 \ln \left( 1 + e^{-\epsilon_k/T} \right) + \text{const}. \end{aligned} \quad (151)$$

**Exercise 4:** Prove via contour integration the following result for the summation over fermionic Matsubara frequencies,

$$T \sum_{k_0} \frac{(k_0 + \xi_1)(k_0 + q_0 + \xi_2)}{(k_0^2 - \epsilon_1^2)[(k_0 + q_0)^2 - \epsilon_2^2]} = -\frac{1}{4\epsilon_1\epsilon_2} \sum_{\epsilon_1, \epsilon_2 = \pm} \frac{(\epsilon_1 - e_1\xi_1)(\epsilon_2 - e_2\xi_2)}{q_0 - e_1\epsilon_1 + e_2\epsilon_2} \frac{f_F(-e_1\epsilon_1)f_F(e_2\epsilon_2)}{f_B(-e_1\epsilon_1 + e_2\epsilon_2)}, \quad (152)$$

where  $k_0 = -i\omega_n$  with fermionic Matsubara frequencies  $\omega_n$ , and  $q_0 = -i\omega_m$  with bosonic (!) Matsubara frequencies  $\omega_m$ , and where  $\xi_1, \xi_2, \epsilon_1, \epsilon_2 > 0$  are real numbers.

### E. Thermodynamic potential for fermions

The result for the Matsubara sum (151) can now be inserted into the partition function (145) to obtain

$$\ln Z = 2V \int \frac{d^3k}{(2\pi)^3} \left[ \frac{\epsilon_k}{T} + \ln \left( 1 + e^{-(\epsilon_k - \mu)/T} \right) + \ln \left( 1 + e^{-(\epsilon_k + \mu)/T} \right) \right]. \quad (153)$$

Consequently, the thermodynamic potential  $\Omega = -T \ln Z$  becomes

$$\frac{\Omega}{V} = -2 \int \frac{d^3k}{(2\pi)^3} \left[ \epsilon_k + T \ln \left( 1 + e^{-(\epsilon_k - \mu)/T} \right) + T \ln \left( 1 + e^{-(\epsilon_k + \mu)/T} \right) \right]. \quad (154)$$

Note the overall factor 2 which accounts for the two spin states of the spin-1/2 fermion. Together with the particle/antiparticle degree of freedom (from  $\mu = \pm 1$ ) we thus see all four degrees of freedom of the Dirac spinor.

[End of 6th lecture, Nov 10th, 2008]

## VII. GAUGE FIELDS

### A. Lagrangians for QCD and QED

In this section we shall compute the partition function for gauge fields. Many applications of thermal field theory in modern research can be found in Quantum Chromodynamics (QCD), for instance heavy-ion collisions and neutron star (quark star) physics. We shall, for the calculation of the partition function, focus on the simpler case of Quantum Electrodynamics (QED). But first we write down the QCD Lagrangian from which we obtain the QED Lagrangian as a limit. We have

$$\mathcal{L}_{\text{QCD}} = -\frac{1}{2} G_{\mu\nu} G^{\mu\nu} + \bar{\psi} (i\gamma^\mu D_\mu + \gamma^0 \mu - m) \psi. \quad (155)$$

Let us explain the meaning of the various quantities and their structure. The field strengths are

$$G_{\mu\nu} = \partial_\mu A_\nu - \partial_\nu A_\mu - ig[A_\mu, A_\nu], \quad (156)$$

where  $g$  is the QCD coupling constant, and where  $A_\mu$  are matrices in the Lie Algebra of the gauge group  $SU(N_c)$  where  $N_c = 3$  is the number of colors. Here,  $SU(N_c)$  is the group of unitary  $N_c \times N_c$  matrices with determinant 1. The dimension of  $SU(N_c)$  is  $N_c^2 - 1$ , thus in this case there are eight generators  $T_a$  which fulfil

$$[T_a, T_b] = if_{abc}T_c, \quad T_a^\dagger = T_a, \quad \text{Tr}[T_a T_b] = \frac{\delta_{ab}}{2}. \quad (157)$$

The generators (more precisely, twice the generators  $\lambda_a = 2T_a$ ) are called Gell-Mann matrices. The gauge fields, which are called gluons, and field strengths can thus be written as

$$A_\mu = A_\mu^a T_a, \quad G_{\mu\nu} = G_{\mu\nu}^a T_a, \quad G_{\mu\nu}^a = \partial_\mu A_\nu^a - \partial_\nu A_\mu^a + gf^{abc}A_\mu^b A_\nu^c. \quad (158)$$

The Dirac spinors  $\psi$  describe quarks and are spinors in a  $4N_f N_c$ -dimensional space with the number of flavors  $N_f$ ; the covariant derivative is

$$D_\mu = \partial_\mu - igA_\mu. \quad (159)$$

With fundamental color indices  $\alpha, \beta \leq 3$ , the adjoint color index  $a \leq 8$ , and flavor indices  $i, j \leq N_f$  we can thus write the Lagrangian as

$$\mathcal{L}_{\text{QCD}} = -\frac{1}{4}G_{\mu\nu}^a G_{\mu\nu}^{a\mu\nu} + \bar{\psi}_i^\alpha \delta_{ij} [i\gamma^\mu (\delta^{\alpha\beta} \partial_\mu - igA_\mu^a T_a^{\alpha\beta}) + \delta^{\alpha\beta} (\gamma^0 \mu_i - m_i)] \psi_j^\beta. \quad (160)$$

Here  $m$  and  $\mu$  are matrices in flavor space, with different masses and chemical potentials for different flavors.

The Lagrangian is invariant under gauge transformations  $U = e^{ig\theta_a(X)T^a} \in SU(N_c)$ . The fermion fields and the gauge fields transform as

$$\psi \rightarrow U\psi, \quad A_\mu \rightarrow UA_\mu U^{-1} + \frac{i}{g}U\partial_\mu U^{-1}, \quad (161)$$

where  $U = U(\mathbf{x}, t)$  may depend on space-time, i.e., the symmetry is local. We can easily check that the Lagrangian is invariant under gauge transformations: one uses  $0 = \partial_\mu(UU^{-1}) = (\partial_\mu U)U^{-1} + U(\partial_\mu U^{-1})$  to find

$$G_{\mu\nu} \rightarrow UG_{\mu\nu}U^{-1}. \quad (162)$$

Then, since  $G_{\mu\nu}G^{\mu\nu}$  is trivial in color space, we see that this term is invariant under gauge transformations. For the quark part we find

$$D_\mu \psi \rightarrow UD_\mu \psi, \quad (163)$$

from which we conclude that  $\bar{\psi}D_\mu \psi$  is invariant and thus we see that  $\mathcal{L}_{\text{QCD}}$  is invariant.

For simplicity, we shall consider QED in the following calculation. In this case the gauge group is  $U(1)$  which is an abelian symmetry. For many physical applications and many calculations this makes the theory tremendously simpler than QCD. For the latter, controlled rigorous calculations from first principles are only valid for very few systems such as systems at very large densities or temperatures. This is due to asymptotic freedom which makes the theory weakly coupled for large momentum transfers. In many other cases, however, the theory is strongly coupled and the theoretical treatment becomes very complicated.

In QED there is no commutator term in the field strengths,

$$F_{\mu\nu} = \partial_\mu A_\nu - \partial_\nu A_\mu, \quad (164)$$

and a gauge transformation is simply given by

$$U(X) = e^{ie\alpha(X)}, \quad A_\mu \rightarrow A_\mu + \frac{i}{e}U\partial_\mu U^{-1} = A_\mu - \partial_\mu \alpha. \quad (165)$$

Since  $U(1)$  is a one-dimensional Lie group, there is only one gauge boson, the photon (compared to eight gluons in QCD). Due to the missing commutator term, the photon has no self-coupling (whereas gluons interact with each other). The fermions are leptons instead of quarks, and the coupling is denoted by  $e$  instead of  $g$ . The Lagrangian, invariant under  $U(1)$ , is

$$\mathcal{L}_{\text{QED}} = -\frac{1}{4}F_{\mu\nu}F^{\mu\nu} + \bar{\psi}(i\gamma^\mu D_\mu + \gamma^0 \mu - m)\psi, \quad (166)$$

with the covariant derivative

$$D_\mu = \partial_\mu - ieA_\mu. \quad (167)$$

### B. Partition function in QED

We now focus on the gauge part of the QED Lagrangian (166), i.e., we are interested in

$$\mathcal{L} = -\frac{1}{4}F_{\mu\nu}F^{\mu\nu} = \frac{1}{2}F_{0i}F_{0i} - \frac{1}{4}F_{ij}F_{ij}. \quad (168)$$

The electric and magnetic fields are given by

$$E_i = -F_{0i} = F_{i0}, \quad \mathbf{B} = \nabla \times \mathbf{A} \Rightarrow B_i = \frac{1}{2}\epsilon_{ijk}F_{jk}. \quad (169)$$

We thus have

$$B^2 = \frac{1}{2}F_{jk}F_{jk}, \quad (170)$$

and the Lagrangian becomes

$$\mathcal{L} = \frac{1}{2}E^2 - \frac{1}{2}B^2. \quad (171)$$

In the following we shall work in the so-called axial gauge

$$A_3 = 0. \quad (172)$$

This does not completely fix the gauge and we will see how the residual gauge freedom appears. The conjugate momenta are

$$\pi_\mu = \frac{\partial \mathcal{L}}{\partial(\partial_0 A_\mu)} = (0, F_{0i}) = F_{0\mu}. \quad (173)$$

We see that there is no momentum conjugate to  $A_0$ . Consequently,  $A_0$  is not a dynamical field. Formally, there is a conjugate momentum

$$\pi_3 = -E_3. \quad (174)$$

However,  $A_3 = 0$  in the chosen gauge, i.e.,  $\pi_3$  is not an independent variable. It can be determined from Gauss' law, which, in the absence of charges, is

$$\nabla \cdot \mathbf{E} = 0. \quad (175)$$

Consequently, we have  $\partial_3 E_3 = \partial_1 \pi_1 + \partial_2 \pi_2$  and thus

$$E_3 = \int_{x_{30}}^{x_3} dx'_3 (\partial_1 \pi_1 + \partial_2 \pi_2) + P(x_1, x_2, t), \quad (176)$$

and

$$A_0 = \int_{x_{30}}^{x_3} dx'_3 E_3 + Q(x_1, x_2, t). \quad (177)$$

The integration constants  $P$  and  $Q$  correspond to the residual gauge freedom. Next we determine the Hamiltonian in terms of the independent variables  $\pi_1, \pi_2, A_1, A_2$ ,

$$\mathcal{H} = \pi_1 \partial_0 A_1 + \pi_2 \partial_0 A_2 - \mathcal{L} \quad (178)$$

We use  $\partial_0 A_i = \pi_i + \partial_i A_0$  (from Eq. (173)) and Eq. (170) to obtain

$$\begin{aligned} \mathcal{H} &= \frac{1}{2}(\pi_1^2 + \pi_2^2) - \frac{1}{2}E_3^2 + \frac{1}{2}B^2 + \pi_1 \partial_1 A_0 + \pi_2 \partial_2 A_0 \\ &= \frac{1}{2}(\pi_1^2 + \pi_2^2) + \frac{1}{2}E_3^2 + \frac{1}{2}B^2, \end{aligned} \quad (179)$$

where we used partial integration and dropped the surface terms (i.e., this identity only holds under the integral  $d^3x$ ). The Hamiltonian now has the familiar form  $\mathcal{H} = E^2/2 + B^2/2$ . The partition function for the bosonic fields  $A_1$ ,  $A_2$  and their conjugate momenta is

$$Z = \int \mathcal{D}\pi_1 \mathcal{D}\pi_2 \int_{\text{periodic}} \mathcal{D}A_1 \mathcal{D}A_2 \exp \int_X (i\pi_1 \partial_\tau A_1 + i\pi_2 \partial_\tau A_2 - \mathcal{H}) . \quad (180)$$

We rewrite the partition function in the following way. First we insert

$$1 = \int \mathcal{D}\pi_3 \delta(\pi_3 + E_3(\pi_1, \pi_2)) . \quad (181)$$

This can be rewritten upon using

$$\delta(\nabla \cdot \boldsymbol{\pi}) = \left( \det \frac{\partial(\nabla \cdot \boldsymbol{\pi})}{\partial\pi_3} \right)^{-1} \delta(\pi_3 + E_3(\pi_1, \pi_2)) . \quad (182)$$

Here one should remember the more familiar form of this identity

$$\delta(f(x)) = \frac{1}{|f'(x_0)|} \delta(x - x_0) , \quad (183)$$

where  $x_0$  is the zero of the function  $f$ . Moreover we use

$$\det \frac{\partial(\nabla \cdot \boldsymbol{\pi})}{\partial\pi_3} = \det(\partial_3) , \quad (184)$$

and we write the  $\delta$ -function in its integral representation,

$$\delta(\nabla \cdot \boldsymbol{\pi}) = \int \mathcal{D}A_0 \exp \left( i \int_X A_0 \nabla \cdot \boldsymbol{\pi} \right) . \quad (185)$$

Here, in the exponential, we have replaced  $A_0 \rightarrow iA_0$  since this yields the replacement  $i \int d^4x A_0 \nabla \cdot \boldsymbol{\pi} \rightarrow i \int_X A_0 \nabla \cdot \boldsymbol{\pi}$  (note that we also have to replace  $dx_0$  by  $-id\tau$ )<sup>1</sup>. Inserting all this into Eq. (181) yields

$$1 = \int \mathcal{D}\pi_3 \int \mathcal{D}A_0 \det(\partial_3) \exp \left( i \int_X A_0 \nabla \cdot \boldsymbol{\pi} \right) , \quad (186)$$

and the partition function becomes (after a partial integration  $A_0 \nabla \cdot \boldsymbol{\pi} \rightarrow -(\nabla A_0) \cdot \boldsymbol{\pi}$ )

$$Z = \int \mathcal{D}\pi_1 \mathcal{D}\pi_2 \mathcal{D}\pi_3 \int_{\text{periodic}} \mathcal{D}A_0 \mathcal{D}A_1 \mathcal{D}A_2 \det(\partial_3) \exp \int_X \left[ i\pi_1 \partial_\tau A_1 + i\pi_2 \partial_\tau A_2 - i(\nabla A_0) \cdot \boldsymbol{\pi} - \frac{1}{2}\pi^2 - \frac{1}{2}B^2 \right] . \quad (187)$$

[End of 7th lecture, Nov 17th, 2008]

The momentum integral now becomes trivial as we have seen in the case of scalar bosons. To this end, we rewrite the exponential with the help of

$$i\pi_1 \partial_\tau A_1 + i\pi_2 \partial_\tau A_2 - i(\nabla A_0) \cdot \boldsymbol{\pi} - \frac{1}{2}\pi^2 = -\frac{1}{2}(\boldsymbol{\pi} - i\partial_\tau \mathbf{A} + i\nabla A_0)^2 - \frac{1}{2}(\partial_\tau \mathbf{A} - \nabla A_0)^2 , \quad (188)$$

where  $\mathbf{A} = (A_1, A_2, 0)$  in the axial gauge we use. Now the integration over the shifted momentum  $\boldsymbol{\pi} - i\partial_\tau \mathbf{A} + i\nabla A_0$  can be performed and yields an irrelevant constant factor which we omit in the following. Consequently,

$$Z = \int_{\text{periodic}} \mathcal{D}A_0 \mathcal{D}A_1 \mathcal{D}A_2 \det(\partial_3) \exp \int_X \mathcal{L} . \quad (189)$$

---

<sup>1</sup> Another way of saying this is that in the field strength  $F_{i0} = \partial_i A_0 - \partial_0 A_i$  for finite temperature we have to replace  $\partial_0$  by  $i\partial_\tau$ . To get the same factor  $i$  from the first term we need to replace  $A_0$  by  $iA_0$ .

We have recovered the Lagrangian in the exponential since

$$-(\partial_\tau \mathbf{A} - \nabla A_0)^2 = E^2. \quad (190)$$

(To see this, one simply ‘‘undoes’’ the finite-temperature replacements  $\partial_0 \rightarrow i\partial_\tau$ ,  $A_0 \rightarrow iA_0$ .) Hence we get the Lagrangian in the form (171).

Before we proceed with Eq. (189) we notice that the general form of the partition function, without specifying a gauge, is

$$Z = \int_{\text{periodic}} \mathcal{D}A_\mu \delta(F) \det \frac{\partial F}{\partial \alpha} \exp \int_X \mathcal{L}, \quad (191)$$

where  $\mathcal{D}A_\mu \equiv \mathcal{D}A_0 \mathcal{D}A_1 \mathcal{D}A_2 \mathcal{D}A_3$ , where  $F$  is a function of the gauge fields and the condition  $F = 0$  fixes the gauge. In our case,  $F = A'_3 = A_3 - \partial_3 \alpha$ . Then, with  $\partial F / \partial \alpha = \partial_3$  we recover Eq. (189). The more general form shows that we integrate over the space of gauge fields ‘‘modulo gauge transformations’’. In other words, for each point in the space of gauge fields, we choose a fixed gauge given by the function  $F$  and fixed by the factor  $\delta(F)$ . Then  $\det(\partial F / \partial \alpha)$  is the determinant of the Jacobian of the transformation  $A'_\mu = F(A_\mu) = A_\mu - \partial_\mu \alpha$ , i.e., it accounts for the change of integration variables according to the gauge transformation. The partition function in the form (191) is manifestly gauge invariant.

Let us now come back to our expression (189) in the axial gauge and compute the functional integral. With Eq. (170) we find

$$\begin{aligned} E^2 - B^2 &= -(\partial_\tau \mathbf{A} - \nabla A_0)^2 - B^2 \\ &= -(\partial_\tau \mathbf{A})^2 - (\nabla A_0)^2 + 2\partial_\tau \mathbf{A} \cdot \nabla A_0 \\ &\quad - (\partial_1 A_2)^2 - (\partial_2 A_1)^2 - (\partial_3 A_1)^2 - (\partial_3 A_2)^2 + 2(\partial_1 A_2)(\partial_2 A_1). \end{aligned} \quad (192)$$

As above, we introduce the Fourier transform of the gauge fields,

$$A_\mu(X) = \frac{1}{\sqrt{TV}} \sum_K e^{-iK \cdot X} A_\mu(K). \quad (193)$$

This yields

$$\int_X (\partial_\tau \mathbf{A})^2 = -\frac{1}{T^2} \sum_K k_0^2 \mathbf{A}(-K) \cdot \mathbf{A}(K), \quad (194a)$$

$$\int_X (\nabla A_0)^2 = \frac{1}{T^2} \sum_K k^2 A_0(-K) A_0(K), \quad (194b)$$

$$\int_X \partial_\tau \mathbf{A} \cdot \nabla A_0 = \frac{1}{T^2} \sum_K ik_0 \mathbf{k} \cdot \mathbf{A}(-K) A_0(K) = \frac{1}{T^2} \sum_K ik_0 \mathbf{k} \cdot \mathbf{A}(K) A_0(-K), \quad (194c)$$

$$\int_X (\partial_1 A_2)^2 = \frac{1}{T^2} \sum_K k_1^2 A_2(-K) A_2(K), \quad (194d)$$

$$\int_X (\partial_1 A_2)(\partial_2 A_1) = \frac{1}{T^2} \sum_K k_1 k_2 A_1(-K) A_2(K) = \frac{1}{T^2} \sum_K k_1 k_2 A_1(K) A_2(-K). \quad (194e)$$

The other terms  $(\partial_2 A_1)^2$ ,  $(\partial_3 A_1)^2$ ,  $(\partial_3 A_2)^2$  are obtained analogously to Eq. (194d). We thus find

$$\int_X \mathcal{L} = -\frac{1}{2T^2} \sum_K (A_0(-K), A_1(-K), A_2(-K)) \begin{pmatrix} k^2 & -ik_0 k_1 & -ik_0 k_2 \\ -ik_0 k_1 & -k_0^2 + k_2^2 + k_3^2 & -k_1 k_2 \\ -ik_0 k_2 & -k_1 k_2 & -k_0^2 + k_1^2 + k_3^2 \end{pmatrix} \begin{pmatrix} A_0(K) \\ A_1(K) \\ A_2(K) \end{pmatrix}. \quad (195)$$

The  $3 \times 3$  matrix is the inverse gauge field propagator in momentum space which we denote by  $\mathcal{D}_0^{-1}(K)$ . Here we have symmetrized the appearing matrix in the exponential. This is important since  $\mathbf{A}(K)$  and  $\mathbf{A}(-K)$  are not independent variables. So suppose we had used some asymmetric ‘‘propagator’’  $\tilde{\mathcal{D}}_0$ . Then we have to write

$$\begin{aligned} \sum_K A_a(-K) [\tilde{\mathcal{D}}_0^{-1}(K)]_{ab} A_b(K) &= \sum_{K>0} A_a(-K) [\tilde{\mathcal{D}}_0^{-1}(K)]_{ab} A_b(K) + \sum_{K<0} A_a(-K) [\tilde{\mathcal{D}}_0^{-1}(K)]_{ab} A_b(K) \\ &= \sum_{K>0} A_a(-K) \left\{ \tilde{\mathcal{D}}_0^{-1}(K)_{ab} + \tilde{\mathcal{D}}_0^{-1}(K)_{ba} \right\} A_b(K), \end{aligned} \quad (196)$$



and arrive at the symmetrized propagator.

We can now use Eq. (57) for the integration to obtain

$$\begin{aligned}
Z &= \det(\partial_3) \left( \det \frac{\mathcal{D}_0^{-1}}{T^2} \right)^{-1/2} \\
&= \det(\partial_3) \left( \prod_K \frac{K^4 k_3^2}{T^6} \right)^{-1/2} \\
&= \det(\partial_3) \left( \prod_K \frac{K^2}{T^2} \right)^{-2/2} \left( \prod_K \frac{k_3^2}{T^2} \right)^{-1/2},
\end{aligned} \tag{197}$$

and thus

$$\ln Z = \ln \det(\partial_3) - 2 \frac{1}{2} \sum_K \ln \frac{k_0^2 - k^2}{T^2} - \frac{1}{2} \sum_K \ln \frac{k_3^2}{T^2}. \tag{198}$$

It remains to evaluate the so-called Fadeev-Popov determinant  $\det(\partial_3)$ . With Eq. (140) we can write this determinant as a functional integral over Grassmann variables  $\bar{C}$ ,  $C$ ,

$$\det(\partial_3) = \int \mathcal{D}\bar{C}\mathcal{D}C \exp \left( - \int_X \bar{C} \partial_3 C \right). \tag{199}$$

Here  $C$  is a complex, scalar field, i.e., it seems to describe a spin-0 boson. On the other hand, the integration goes over Grassmann variables, indicating fermionic properties. This unphysical field is called a Fadeev-Popov ghost field. It plays a more important role in non-abelian gauge theories but we see that it is needed also here. With the Fourier transform

$$C(X) = \frac{1}{\sqrt{V}} \sum_K e^{-iK \cdot X} C(K), \tag{200}$$

(bosonic Matsubara frequencies!) we have

$$- \int_X \bar{C} \partial_3 C = - \sum_K \bar{C}(K) \frac{ik_3}{T} C(K). \tag{201}$$

Consequently, the ghost contribution is

$$\det(\partial_3) = \det \frac{ik_3}{T} = \prod_K \frac{ik_3}{T} \propto \prod_K \frac{k_3}{T}. \tag{202}$$

We see that this term exactly cancels the third term on the right-hand side of Eq. (198) and we are left with

$$\ln Z = -2 \frac{1}{2} \sum_K \ln \frac{k_0^2 - k^2}{T^2}. \tag{203}$$

This result shows the two degrees of freedom of the gauge field. The third degree of freedom, unphysical due to gauge symmetry, is cancelled by the ghosts.

## VIII. INTERACTIONS

### A. Perturbative expansion in $\lambda\phi^4$ theory

We add an interaction term with coupling constant  $\lambda$  to the Lagrangian for a real scalar field (42) to obtain the Lagrangian

$$\mathcal{L} = \mathcal{L}_0 + \mathcal{L}_I = \frac{1}{2} \partial_\mu \phi \partial^\mu \phi - \frac{1}{2} m^2 \phi^2 - \lambda \phi^4. \tag{204}$$

We use the index 0 for the contribution we have already computed above. The partition function then is

$$Z = \int \mathcal{D}\phi e^S, \quad (205)$$

with the action

$$S = S_0 + S_I = \int_X \mathcal{L}_0 + \int_X \mathcal{L}_I, \quad S_I = -\lambda \int_X \phi^4. \quad (206)$$

Without interaction,  $S_I = 0$ , we could compute  $\ln Z$  exactly. In the presence of interactions this is not possible. Therefore, we need to apply an approximation. The simplest approximation is to truncate the expansion in the coupling  $\lambda$  under the assumption that the coupling is small. Denoting the noninteracting part by

$$Z_0 \equiv \int \mathcal{D}\phi e^{S_0} \quad (207)$$

we can write the expansion as

$$\begin{aligned} \ln Z &= \ln \int \mathcal{D}\phi e^{S_0 + S_I} \\ &= \ln \int \mathcal{D}\phi e^{S_0} \sum_{n=0}^{\infty} \frac{S_I^n}{n!}. \end{aligned} \quad (208)$$

Now if we add and subtract  $\ln Z_0$  we can write this as

$$\begin{aligned} \ln Z &= \ln Z_0 + \ln \frac{\int \mathcal{D}\phi e^{S_0} \sum_{n=0}^{\infty} \frac{S_I^n}{n!}}{\int \mathcal{D}\phi e^{S_0}} \\ &= \ln Z_0 + \ln Z_I, \end{aligned} \quad (209)$$

with

$$\ln Z_I \equiv \ln \left( 1 + \sum_{n=1}^{\infty} \frac{1}{n!} \frac{\int \mathcal{D}\phi e^{S_0} S_I^n}{\int \mathcal{D}\phi e^{S_0}} \right) = \ln \left( 1 + \sum_{n=1}^{\infty} \frac{\langle S_I^n \rangle_0}{n!} \right). \quad (210)$$

Here  $\langle - \rangle_0$  denotes the ensemble average over the noninteracting ensemble. From the definition of  $S_I$  we know that each factor of  $S_I$  comes with one power of  $\lambda$ . If we expand  $\ln Z_I$  to, say, third order in the coupling, we thus obtain, using  $\ln(1+x) = \sum_{n=1}^{\infty} (-1)^{n+1} x^n/n$ ,

$$\begin{aligned} \ln Z_I &\simeq \ln \left( 1 + \langle S_I \rangle_0 + \frac{\langle S_I^2 \rangle_0}{2} + \frac{\langle S_I^3 \rangle_0}{6} \right) \\ &\simeq \langle S_I \rangle_0 + \frac{1}{2} (\langle S_I^2 \rangle_0 - \langle S_I \rangle_0^2) + \frac{1}{6} (\langle S_I^3 \rangle_0 - 3\langle S_I \rangle_0 \langle S_I^2 \rangle_0 + 2\langle S_I \rangle_0^3), \end{aligned} \quad (211)$$

where we have ordered the contributions according to the powers  $\lambda$ ,  $\lambda^2$ ,  $\lambda^3$ .

[End of 8th lecture, Nov 24th, 2008]

Let us compute the first correction  $\propto \lambda$  explicitly. We have

$$\langle S_I \rangle_0 = -\lambda \frac{\int \mathcal{D}\phi e^{S_0} \int_X \phi^4(X)}{\int \mathcal{D}\phi e^{S_0}}. \quad (212)$$

From Sec. IV we know that

$$e^{S_0} = \exp \left[ -\frac{1}{2} \sum_K \phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K) \right] = \prod_K \exp \left[ -\frac{1}{2} \phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K) \right], \quad (213)$$

with the inverse propagator

$$D_0^{-1}(K) = \omega_n^2 + k^2 + m^2. \quad (214)$$

In momentum space, the  $\phi^4$  term becomes

$$\begin{aligned} \int_X \phi^4(X) &= \frac{1}{T^2 V^2} \sum_{K_1, \dots, K_4} \int_X e^{i(K_1 + \dots + K_4) \cdot X} \phi(K_1) \dots \phi(K_4) \\ &= \frac{1}{T^3 V} \sum_{K_1, \dots, K_4} \delta(K_1 + \dots + K_4) \phi(K_1) \dots \phi(K_4). \end{aligned} \quad (215)$$

Inserting Eqs. (213) and (215) into Eq. (212) yields

$$\langle S_I \rangle_0 = -\frac{\lambda}{T^3 V} \frac{\sum_{K_1, \dots, K_4} \delta(K_1 + \dots + K_4) \prod_K \int d\phi(K) e^{-\frac{1}{2}\phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K)} \phi(K_1) \dots \phi(K_4)}{\prod_K \int d\phi(K) e^{-\frac{1}{2}\phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K)}}. \quad (216)$$

The integral in the numerator is only nonvanishing if the four momenta  $K_1, K_2, K_3, K_4$  cancel each other pairwise. Otherwise, if there is a single power of  $\phi(K)$ , the integral over  $\phi(K)$  is zero by symmetry. (Remember that  $\phi(-K) = \phi^*(K)$  and thus  $\phi(K)\phi(-K) = |\phi(K)|^2$ .) Hence we have for instance  $K_1 = -K_2 \equiv Q$  and  $K_3 = -K_4 \equiv P$ , and the delta-function is automatically fulfilled. There are 3 possibilities for the momenta to be pairwise identical and thus we obtain

$$\langle S_I \rangle_0 = -\frac{3\lambda}{T^3 V} \sum_{Q, P} \frac{\prod_K \int d\phi(K) e^{-\frac{1}{2}\phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K)} \phi(-Q) \phi(Q) \phi(-P) \phi(P)}{\prod_K \int d\phi(K) e^{-\frac{1}{2}\phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K)}}. \quad (217)$$

Now we notice that all integrals over  $K \neq P, Q$  appear identically in numerator and denominator and thus cancel. The  $\phi(P)$  and  $\phi(Q)$  integrals factorize and we obtain

$$\langle S_I \rangle_0 = -\frac{3\lambda}{T^3 V} \left[ \sum_Q \frac{\int d\phi(Q) e^{-\frac{1}{2}\phi(-Q) \frac{D_0^{-1}(Q)}{T^2} \phi(Q)} \phi(Q) \phi(-Q)}{\int d\phi(Q) e^{-\frac{1}{2}\phi(-Q) \frac{D_0^{-1}(Q)}{T^2} \phi(Q)}} \right]^2. \quad (218)$$

Now we use

$$\frac{\int_{-\infty}^{\infty} dx x^2 e^{-ax^2/2}}{\int_{-\infty}^{\infty} dx e^{-ax^2/2}} = \frac{1}{a}, \quad (219)$$

to obtain

$$\ln Z_I^{(1)} = \langle S_I \rangle_0 = -3\lambda \frac{T}{V} \left[ \sum_Q D_0(Q) \right]^2. \quad (220)$$

We shall compute this expression in Sec. VIII C. Here we proceed by introducing Feynman diagrams: it is convenient to translate the complicated algebraic perturbative expansion into a diagrammatic form. One starts by representing each field by a line with a direction, the direction indicating whether the field is ingoing or outgoing (i.e., inverting the direction corresponds to  $\phi(K) \rightarrow \phi(-K)$ ). Then the interaction term  $-\lambda\phi^4$  is represented in momentum space by



$$(221)$$

By convention, we have chosen the signs of the momenta such that all lines are ingoing. Since the momenta  $K_1, K_2, K_3, K_4$  have to cancel pairwise, we connect the lines pairwise and interpret each resulting line as a propagator. (From the explicit integration above we know how the propagators come about.) There are three possibilities to connect the four lines pairwise and thus the algebraic result is translated into a Feynman diagram as follows,

$$\langle S_I \rangle_0 = -3\lambda \frac{T}{V} \left[ \sum_Q D_0(Q) \right]^2 = 3 \text{ (diagram of two adjacent circles)} \quad (222)$$

In summary, the vertex gives a factor  $-\lambda$ , the factor 3 is a combinatorial factor, the closed line is a propagator  $T/V \sum_Q D_0(Q)$ , and momentum conservation gives  $V/T \delta(K_{\text{in}} - K_{\text{out}})$  which here is automatically fulfilled and thus simply gives a factor  $V/T$ . In general, the rules to find all contributions to the partition function in a given order  $\lambda^n$  are

1. Draw all connected diagrams with combinatorial prefactors.
2. Each (closed) line gives a propagator  $\frac{T}{V} \sum_K D_0(K)$ .
3. Each vertex gives a factor  $-\lambda$  and a momentum-conserving Kronecker-delta  $\frac{V}{T} \delta(K_{\text{in}} - K_{\text{out}})$ .

In the first-order contribution it is clear that there is only a connected diagram. We shall explain now, for the second-order corrections to  $\ln Z$ , why the disconnected diagrams cancel. The second-order terms of the partition function are

$$\ln Z_I^{(2)} = \frac{1}{2} (\langle S_I^2 \rangle_0 - \langle S_I \rangle_0^2). \quad (223)$$

From Eq. (222) we know the diagrammatic representation of  $\langle S_I \rangle_0^2$ . For  $\langle S_I^2 \rangle_0$  we need to start from  $(-\lambda\phi^4)^2$  which, in analogy to Eq. (221) is represented as

$$\text{(diagram of two crossed lines with momenta } K_1, K_2, K_3, K_4 \text{ and } K_5, K_6, K_7, K_8 \text{)} \quad (224)$$

Again we have to construct all possible diagrams by connecting the eight lines pairwise. One of the diagrams we obtain is the product of two disconnected “double-bubbles” (each with a combinatorial factor 3) which exactly cancels the term  $\langle S_I \rangle_0^2$ . We are left with

$$\ln Z_I^{(2)} = 36 \text{ (diagram of three adjacent circles)} + 12 \text{ (diagram of two overlapping circles)} \quad (225)$$

The fact that the disconnected diagrams cancel out is general, i.e., Eq. (210) simplifies to

$$\ln Z_I = \sum_{n=1}^{\infty} \frac{\langle S_I^n \rangle_{0,\text{connected}}}{n!}. \quad (226)$$

Finally, let us translate the second-order diagrams back into momentum sums:

$$\begin{aligned} \text{(diagram of three adjacent circles)} &= (-\lambda)^2 \left(\frac{T}{V}\right)^4 \left(\frac{V}{T}\right)^2 \sum_{K_1, \dots, K_4} \delta(K_2 + K_3) D_0(K_1) \dots D_0(K_4) \\ &= \lambda^2 \frac{T^2}{V^2} \left[ \sum_K D_0(K) \right]^2 \sum_{K_2, K_3} \delta(K_2 + K_3) D_0(K_2) D_0(K_3), \end{aligned} \quad (227a)$$

$$\text{(diagram of two overlapping circles)} = (-\lambda)^2 \left(\frac{T}{V}\right)^4 \left(\frac{V}{T}\right)^2 \sum_{K_1, \dots, K_4} \delta(K_1 + \dots + K_4) D_0(K_1) \dots D_0(K_4). \quad (227b)$$

### B. Propagator, self-energy, and one-particle irreducible (1PI) diagrams

From above, see Eq. (219), we know that the free propagator can be written as an ensemble average over the noninteracting ensemble,

$$\begin{aligned} D_0(Q) &= \frac{1}{T^2} \frac{\prod_K \int d\phi(K) e^{-\frac{1}{2}\phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K)} \phi(Q) \phi(-Q)}{\prod_K \int d\phi(K) e^{-\frac{1}{2}\phi(-K) \frac{D_0^{-1}(K)}{T^2} \phi(K)}} \\ &= \frac{1}{T^2} \langle \phi(Q) \phi(-Q) \rangle_0, \end{aligned} \quad (228)$$

The full propagator, which includes interactions to all orders, can then also be written as an ensemble average over the interacting ensemble,

$$D(Q) = \frac{1}{T^2} \langle \phi(Q) \phi(-Q) \rangle. \quad (229)$$

We now show that this is the form of the propagator in momentum space in the case of a translationally invariant system, given that the general propagator in position space is defined as

$$D(X_1, X_2) \equiv \langle \phi(X_1) \phi(X_2) \rangle. \quad (230)$$

If the system is translationally invariant, this propagator only depends on the difference  $X_1 - X_2$ . Without loss of generality we can thus set  $X_2 = 0$  and denote  $X = X_1$ . Then, the Fourier transform is

$$\begin{aligned} D(Q) &= \int_X e^{iQ \cdot X} \langle \phi(X) \phi(0) \rangle \\ &= \frac{1}{TV} \sum_{K_1, K_2} \int_X e^{iQ \cdot X} e^{-iK_1 \cdot X} \langle \phi(K_1) \phi(K_2) \rangle \\ &= \frac{1}{T^2} \sum_{K_2} \langle \phi(Q) \phi(K_2) \rangle \\ &= \frac{1}{T^2} \langle \phi(Q) \phi(-Q) \rangle. \end{aligned} \quad (231)$$

In the last step we have used that the ensemble average is only nonzero for  $K_2 = -Q$ . We have thus arrived at the propagator (229).

**Exercise 5:** Compute the free propagator  $D_0(X, 0)$  in position space and show that it behaves as  $D_0(X, 0) \propto 1/x^2$  for small  $x$ ,  $x \ll 1/T, 1/m$ , and as  $D_0(X, 0) \propto T e^{-mx}/x$  for large  $x$ ,  $x \gg 1/T$  (and  $m \ll T$ ). Here,  $x \equiv |\mathbf{x}|$ . (Hint: In the Matsubara summation, use the function  $1/(e^{\omega/T} - 1)$  instead of  $1/2 \coth \omega/(2T)$  in the analogue of Eq. (63); this ensures that, when closing the contour, the contribution of the infinite semi-circle in the positive half-plane vanishes.)

For a systematic calculation of the perturbation series it is convenient to divide the full (inverse) propagator into a free part and an interaction part, called self-energy  $\Pi$ . We write

$$D^{-1}(K) = D_0^{-1}(K) + \Pi(K). \quad (232)$$

Let us discuss the self-energy up to second order in the coupling. We first observe that the propagator can be written as a functional derivative of  $\ln Z$ . Remember from Eq. (208) that

$$\ln Z = \ln \int \mathcal{D}\phi e^{S_0} e^{S_I}. \quad (233)$$

Therefore,

$$\begin{aligned}
\frac{\delta}{\delta D_0^{-1}} \ln Z &= \frac{1}{\int \mathcal{D}\phi e^{S_0+S_I}} \frac{\delta}{\delta D_0^{-1}} \int \mathcal{D}\phi e^{S_0} e^{S_I} \\
&= \frac{1}{\int \mathcal{D}\phi e^{S_0+S_I}} \frac{\delta}{\delta D_0^{-1}} \prod_K \int d\phi(K) e^{-\frac{1}{2}\phi(-K)\frac{D_0^{-1}}{T^2}\phi(K)} e^{S_I} \\
&= -\frac{1}{2T^2} \frac{\int \mathcal{D}\phi e^{S_0} e^{S_I} \phi(-Q)\phi(Q)}{\int \mathcal{D}\phi e^{S_0+S_I}} \\
&= -\frac{1}{2} D(Q), \tag{234}
\end{aligned}$$

and thus

$$D(Q) = -2 \frac{\delta \ln Z}{\delta D_0^{-1}} = 2D_0^2 \frac{\delta \ln Z}{\delta D_0}. \tag{235}$$

Then, from Eq. (232) we have

$$D = (D_0^{-1} + \Pi)^{-1} = (1 + D_0\Pi)^{-1} D_0. \tag{236}$$

Using Eq. (235) we have

$$(1 + \Pi D_0)^{-1} = D D_0^{-1} = 2D_0 \frac{\delta \ln Z}{\delta D_0}. \tag{237}$$

We now expand the self-energy in powers of  $\lambda$ ,

$$\Pi = \sum_{n=1}^{\infty} \Pi_n, \tag{238}$$

such that  $\Pi_n$  is proportional to  $\lambda^n$ . Then, up to second order in  $\lambda$ , we have

$$(1 + \Pi D_0)^{-1} = 1 - D_0\Pi_1 - D_0\Pi_2 + D_0\Pi_1 D_0\Pi_1 + \mathcal{O}(\lambda^3). \tag{239}$$

This is the left-hand side of Eq. (237). For the right-hand side we use, see Eq. (60),

$$Z_0 = \frac{1}{2} \sum_K \ln(D_0^{-1} T^2) \quad \Rightarrow \quad \frac{\delta \ln Z_0}{\delta D_0} = \frac{1}{2} D_0^{-1}, \tag{240}$$

and thus

$$\begin{aligned}
2D_0 \frac{\delta \ln Z}{\delta D_0} &= 2D_0 \left( \frac{\delta \ln Z_0}{\delta D_0} + \frac{\delta \ln Z_I}{\delta D_0} \right) \\
&= 1 + 2D_0 \frac{\delta \ln Z_I}{\delta D_0} \\
&= 1 + 2D_0 \left[ \frac{\delta \langle S_I \rangle_0}{\delta D_0} + \frac{1}{2} \frac{\delta(\langle S_I^2 \rangle_0 - \langle S_I \rangle_0^2)}{\delta D_0} + \mathcal{O}(\lambda^3) \right]. \tag{241}
\end{aligned}$$

Thus, upon comparing Eqs. (239) and (241) we have

$$\Pi_1 + \Pi_2 - \Pi_1 D_0 \Pi_1 + \dots = -2 \frac{\delta \ln Z_I}{\delta D_0}. \tag{242}$$

The first- and second- order contributions are

$$\Pi_1 = -2 \frac{\delta \langle S_I \rangle_0}{\delta D_0}, \tag{243a}$$

$$\Pi_2 - \Pi_1 D_0 \Pi_1 = -\frac{\delta(\langle S_I^2 \rangle_0 - \langle S_I \rangle_0^2)}{\delta D_0}. \tag{243b}$$

The first-order contribution becomes

$$\begin{aligned}\Pi_1 &= 6\lambda \frac{T}{V} \frac{\delta}{\delta D_0} \left[ \sum_K D_0(K) \right]^2 \\ &= 12\lambda \frac{T}{V} \sum_K D_0(K).\end{aligned}\quad (244)$$

We see that taking the functional derivative with respect to the propagator is equivalent to cutting a line in the Feynman diagram,

$$\begin{aligned}\Pi_1 &= -2 \frac{\delta \langle S_I \rangle_0}{\delta D_0} \\ &= -2 \frac{\delta}{\delta D_0} 3 \text{ (two circles)} \\ &= -12 \text{ (one circle)}\end{aligned}\quad (245)$$

The additional factor of 2 appears since each of the two lines can be cut to obtain the same diagram. With this “cutting rule”, we can easily determine the second-order contributions with the help of the diagrams. By cutting a line in the respective diagrams we obtain

$$\begin{aligned}\Pi_2 - \Pi_1 D_0 \Pi_1 &= -\frac{\delta(\langle S_I^2 \rangle_0 - \langle S_I \rangle_0^2)}{\delta D_0} \\ &= -\frac{\delta}{\delta D_0} \left[ 72 \text{ (three circles)} + 24 \text{ (two circles with internal lines)} \right] \\ &= -144 \text{ (two circles)} - 144 \text{ (two circles)} - 96 \text{ (one circle)}\end{aligned}\quad (246)$$

From Eq. (245) we conclude

$$\Pi_1 D_0 \Pi_1 = 144 \text{ (two circles)} \quad (247)$$

such that we obtain

$$\Pi_2 = -144 \text{ (two circles)} - 96 \text{ (one circle)} \quad (248)$$

We see that the diagram which can be divided into two disconnected diagrams by cutting one line cancels. This is a general fact and the self-energy is given by all diagrams that cannot be divided into two by cutting one line. These diagrams are called “one-particle irreducible (1PI)”, and thus Eq. (242) simplifies to

$$\Pi = -2 \left( \frac{\delta \ln Z_I}{\delta D_0} \right)_{\text{1PI}}. \quad (249)$$

### C. Evaluation of the first-order corrections

We can now compute the first-order contribution to the self-energy (244) and to the pressure from Eq. (220). Making use of Eq. (68) we obtain

$$\Pi_1 = 12\lambda \frac{T}{V} \sum_K \frac{1}{\omega_n^2 + \epsilon_k^2} = \Pi_1^{\text{vac}} + \Pi_1^T, \quad (250)$$

with the temperature-independent vacuum part

$$\Pi_1^{\text{vac}} = \Pi_1(T=0) = 6\lambda \int \frac{d^3k}{(2\pi)^3} \frac{1}{\epsilon_k}, \quad (251)$$

and the temperature-dependent part

$$\Pi_1^T = 12\lambda \int \frac{d^3k}{(2\pi)^3} \frac{f_B(\epsilon_k)}{\epsilon_k}. \quad (252)$$

While  $\Pi_1^T$  is finite, the vacuum part is divergent. This divergence comes from large momenta,  $k \rightarrow \infty$ , and is thus called *ultraviolet* divergence. We can use the usual zero-temperature renormalization to remove this divergence since the finite-temperature part is finite<sup>2</sup>. This is done by adding a counterterm which is chosen such that the renormalized self-energy is obtained by subtracting the vacuum part,

$$\Pi_1^{\text{ren}} = \Pi_1 - \Pi_1(T=0). \quad (253)$$

From Eq. (232) we know that the inverse propagator to first order in  $\lambda$  is

$$D^{-1}(K) = \omega_n^2 + k^2 + m^2 + \Pi_1. \quad (254)$$

This shows that the self-energy plays the role of a mass squared. The counterterm we thus add to the Lagrangian is written as a mass term

$$\mathcal{L} \rightarrow \mathcal{L} - \frac{1}{2} \delta m^2 \phi^2. \quad (255)$$

However, this term is treated as an interaction, i.e.,  $\delta m^2$  has to be thought of as being of order  $\lambda$ . From Eq. (210) we have concluded that the  $\phi^4$  term produces a “double-bubble” diagram for the first-order correction. Analogously, there is a “single-bubble” contribution ( $\delta m^2$  replacing a factor  $\phi^2$ , i.e., one closed loop) from  $\delta m^2 \phi^2$ , denoted as

$$\delta m^2 \langle \phi^2 \rangle_0 = \text{---} \bigcirc \text{---} \quad (256)$$

Its contribution to the self-energy is then obtained from cutting one line,

$$\delta m^2 = \text{---} \times \text{---} \quad (257)$$

This contribution is now chosen such that the condition (253) is fulfilled, i.e.,

$$\delta m^2 = -\Pi_1^{\text{vac}}. \quad (258)$$

After this renormalization we can evaluate the first-order self-energy. For the massless case  $m = 0$  (or, equivalently, for large temperatures  $T \gg m$ ) we can do so analytically,

$$\Pi_1^{\text{ren}} = 12\lambda \int \frac{d^3k}{(2\pi)^3} \frac{f_B(\epsilon_k)}{\epsilon_k} \simeq \lambda T^2, \quad (259)$$

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<sup>2</sup> In the next section we will encounter an *infrared* divergence, more intimately related to finite temperature effects.



where we used

$$\int_0^\infty dx \frac{x}{e^x - 1} = \frac{\pi^2}{6}. \quad (260)$$

We see that a massless boson acquires a *thermal mass*  $\lambda T^2$ . The pressure to first order in the coupling is given by

$$P = \frac{T}{V} \ln Z_0 + \frac{T}{V} \ln Z_I^{(1)}. \quad (261)$$

With the noninteracting partition function from Eq. (73) we have

$$\frac{T}{V} \ln Z_0 = -T \int \frac{d^3k}{(2\pi)^3} \ln(1 - e^{-\epsilon_k/T}) \simeq \frac{\pi^2 T^4}{90}, \quad (262)$$

where we have dropped the  $T$ -independent contribution which is only an irrelevant shift in the pressure, where we have approximated the expression for  $T \gg m$ , and where we have used

$$\int_0^\infty dx x^2 \ln(1 - e^{-x}) = -\frac{\pi^4}{45}. \quad (263)$$

With  $\ln Z_I^{(1)}$  from Eq. (220) plus the contribution from the mass counterterm we have

$$\begin{aligned} \ln Z_I^{(1)} &= 3 \left( \text{two circles} \right) - \frac{1}{2} \left( \text{circle with cross} \right) \\ &= -3\lambda \frac{V}{T} \left[ \frac{T}{V} \sum_Q D_0(Q) \right]^2 - \frac{1}{2} \delta m^2 \sum_Q D_0(Q). \end{aligned} \quad (264)$$

Using the form of the self-energy (244) and the mass counterterm (258) and dividing both terms into vacuum and temperature-dependent parts according to Eqs. (251) and (252) we have

$$\begin{aligned} \ln Z_I^{(1)} &= -\frac{V}{T} \frac{1}{48\lambda} (\Pi_1^{\text{vac}} + \Pi_1^T)^2 + \frac{V}{T} \frac{1}{24\lambda} \Pi_1^{\text{vac}} (\Pi_1^{\text{vac}} + \Pi_1^T) \\ &= \frac{V}{T} \frac{1}{48\lambda} [(\Pi_1^{\text{vac}})^2 - (\Pi_1^T)^2]. \end{aligned} \quad (265)$$

Again, we drop the temperature-independent part to get

$$\frac{T}{V} \ln Z_I^{(1)} = -3\lambda \left[ \int \frac{d^3k}{(2\pi)^3} \frac{f_B(\epsilon_k)}{\epsilon_k} \right]^2 \simeq -\frac{\lambda T^4}{48}, \quad (266)$$

again approximating for  $T \gg m$  and using Eq. (259). Putting Eqs. (262) and (266) together yields the pressure

$$P = T^4 \left( \frac{\pi^2}{90} - \frac{\lambda}{48} + \dots \right). \quad (267)$$

#### D. Infrared divergence and resummation of ring diagrams

We have seen above that the first-order self-energy  $\Pi_1$  gives rise to a thermal mass  $\lambda T^2$ . In particular, if  $m = 0$ , the scalar field acquires a finite mass only through a temperature effect. We shall in the following focus on the case  $m = 0$ . For small energies and momenta, at most of the order of the thermal mass,  $\omega_n^2, k^2 \lesssim \lambda T^2$ , the free inverse propagator  $D_0^{-1} = \omega_n^2 + k^2$  is (at most) of order  $\lambda T^2$ , and, as we have seen, also the correction through the self energy is of the order  $\lambda T^2$ . This indicates that the naive perturbation series might not be the correct procedure. Indeed, we shall see in the following that one needs to “resum” a certain class of infinitely many diagrams because of an *infrared* divergence, i.e., a divergence coming from small momenta (and energies), as the above simple argument suggests.

Before we discuss this resummation more systematically, let us show a very direct form of a resummation, namely to use the full instead of the free propagator in the one-loop expression of the self-energy,

$$\begin{aligned}\Pi &= 12\lambda \frac{T}{V} \sum_K D(K) \\ &= 12\lambda \frac{T}{V} \sum_K \frac{1}{D_0^{-1}(K) + \Pi}.\end{aligned}\tag{268}$$

This is a self-consistent equation for  $\Pi$ . Note that in our simple example of real  $\phi^4$  theory  $\Pi$  does not depend on  $K$ . In other more complicated theories where  $\Pi$  depends on four-momentum, Eq. (268) is a complicated integral equation. But also here the general solution of (268) has to be done numerically. Writing

$$\frac{1}{D_0^{-1}(K) + \Pi} = D_0 \sum_{n=1}^{\infty} (-\Pi D_0)^n,\tag{269}$$

we see that the self energy now is a loop which itself has any arbitrary number of self-energy insertions, which in turn have self-energy insertions and so on. Such a sum, which formally includes all powers of the coupling constant, is usually termed “resummation”. If we replace  $\Pi$  in the denominator of Eq. (268) by the first-order approximation  $\Pi_1$ , the corresponding diagrams consist of a loop with  $n$  loops attached to it, sometimes called “daisy”. Using the full  $\Pi$ , each of the  $n$  loops itself gets additional loops, hence here we sum over “superdaisy” diagrams. (But note that, even if the full self-energy is used, Eq. (268) still has the form of a one-loop self-energy, i.e., even in the “superdaisy” resummation we only sum over a subset of all possible diagrams.)

We evaluate Eq. (268) as follows,

$$\begin{aligned}\Pi &= 12\lambda \int \frac{d^3k}{(2\pi)^3} T \sum_n \frac{1}{\omega_n^2 + k^2 + \Pi} \\ &= 12\lambda \int \frac{d^3k}{(2\pi)^3} \frac{f_B(\sqrt{k^2 + \Pi})}{\sqrt{k^2 + \Pi}},\end{aligned}\tag{270}$$

where we used the Matsubara sum (68) and where we dropped the zero-temperature contribution. With the new integration variable  $x = \sqrt{k^2/\Pi + 1}$  we can write this as

$$1 = \frac{6\lambda}{\pi^2} \int_1^{\infty} dx \sqrt{x^2 - 1} f_B(\Pi^{1/2}x).\tag{271}$$

If one wants to keep all superdaisy diagrams, one has to proceed numerically now.

**Exercise 6:** Solve Eq. (268) numerically and plot  $\Pi^{1/2}/T$  as a function of  $\lambda$ . Compare this curve with the result (259) for  $\Pi_1$ .

We can extract an analytical result by using the expansion

$$\int_1^{\infty} dx \sqrt{x^2 - 1} f_B(ux) = \frac{2\pi^2}{u^2} \left[ \frac{1}{12} - \frac{u}{4\pi} + \mathcal{O}(u^2 \ln u) \right].\tag{272}$$

Inserting this expansion into Eq. (271), solving the resulting equation for  $\Pi$  and expanding in powers of  $\lambda$  yields

$$\Pi = \lambda T^2 - \frac{3T^2 \lambda^{3/2}}{\pi} + \dots\tag{273}$$

Interestingly, besides the first-order term, we have found a term proportional to  $\lambda^{3/2}$ . We shall see now that this power also appears in the thermodynamical potential, and we shall see that it is related to the infrared divergence of certain diagrams.

Let us start from Eq. (249) and separate the first-order contribution which we already have computed,

$$\ln Z_I = \langle S_I \rangle_0 + \sum_{N=2}^{\infty} \frac{\langle S_I^N \rangle_{0,\text{connected}}}{N!}.\tag{274}$$

The second-order diagrams are given in Eqs. (227). We see that the first of these diagrams, Eq. (227a) can be written as

$$\begin{array}{c} \text{Diagram: Three circles in a row, each with an arrow pointing to the right, connected by lines.} \end{array} = \frac{\Pi_1^2}{12^2} V \sum_n \int \frac{d^3 k}{(2\pi)^3} \frac{1}{(\omega_n^2 + k^2)^2}. \quad (275)$$

For  $n = 0$  the integrand goes like  $1/k^2$  which is infrared divergent. One can check that the second diagram, given in Eq. (227b) does not have this property. The solution to this apparent problem is to resum all diagrams of this “dangerous” kind as we did above for the self energy. More precisely, these are the “daisy” diagrams, where  $N$  loops are attached to an interior loop (sometimes also called “ring” diagrams). We have

$$\begin{aligned} \sum_{N=2}^{\infty} \frac{\langle S_I^N \rangle_{0, \text{connected, daisy}}}{N!} &= \sum_{N=2}^{\infty} \frac{1}{N!} 6^N 2^{N-1} \frac{N!}{N} \begin{array}{c} \text{Diagram: A central circle with N smaller circles attached to its circumference.} \\ (N \text{ loops}) \end{array} \\ &= \sum_{N=2}^{\infty} \frac{1}{N!} 6^N 2^{N-1} \frac{N!}{N} V \sum_n \int \frac{d^3 k}{(2\pi)^3} \frac{(-\Pi_1)^N}{12^N} D_0(K)^N \\ &= \frac{V}{2} \sum_n \int \frac{d^3 k}{(2\pi)^3} \sum_{N=2}^{\infty} \frac{1}{N} [-\Pi_1 D_0(K)]^N \\ &= -\frac{V}{2} \sum_n \int \frac{d^3 k}{(2\pi)^3} \{ \ln [1 + \Pi_1 D_0(K)] - \Pi_1 D_0(K) \} \\ &= -\frac{V}{2} \sum_n \int \frac{d^3 k}{(2\pi)^3} \left[ \ln \left( 1 + \frac{\lambda T^2}{\omega_n^2 + k^2} \right) - \frac{\lambda T^2}{\omega_n^2 + k^2} \right] \end{aligned} \quad (276)$$

The origin of the combinatorial factors is:  $6^N$  for choosing one pair of lines from each of the  $N$  crosses;  $2^{N-1}$  for the number of ways to connect the chosen pairs of lines to obtain a ring diagram;  $N!/N$  for the number of ways to order the  $N$  loops around the ring. Let us again look at the zero Matsubara mode,  $n = 0$ . All other Matsubara modes give contributions of higher order in  $\lambda$  (remember that naively, i.e., ignoring any infrared divergence, one would expect every single diagram of the sum to be of order  $\lambda^2$  or higher). With the noninteracting and first-order results (262) and (266) we thus find the pressure

$$P \simeq \frac{\pi^2 T^4}{90} - \frac{\lambda T^4}{48} - \frac{T}{2} \int \frac{d^3 k}{(2\pi)^3} \left[ \ln \left( 1 + \frac{\lambda T^2}{k^2} \right) - \frac{\lambda T^2}{k^2} \right]. \quad (277)$$

With

$$\int dk k^2 \left[ \ln \left( 1 + \frac{a^2}{k^2} \right) - \frac{a^2}{k^2} \right] = -\frac{2a^3}{3} \arctan \frac{k}{a} - \frac{a^2 k}{3} + \frac{k^3}{3} \ln \left( 1 + \frac{a^2}{k^2} \right) \quad (278)$$

we have

$$\int \frac{d^3 k}{(2\pi)^3} \left[ \ln \left( 1 + \frac{\lambda T^2}{k^2} \right) - \frac{\lambda T^2}{k^2} \right] = -\frac{\lambda^{3/2} T^3}{6\pi}, \quad (279)$$

and thus

$$P = \frac{\pi^2 T^4}{90} \left[ 1 - \frac{15}{8} \frac{\lambda}{\pi^2} + \frac{15}{2} \left( \frac{\lambda}{\pi^2} \right)^{3/2} + \dots \right]. \quad (280)$$

Consequently, we have found that the next term in the perturbation series of the pressure is not of order  $\lambda^2$ , but of order  $\lambda^{3/2}$ , as we have seen above for the self-energy.

**Exercise 7:** Derive the lowest-order correction  $\langle S_I^2 \rangle_0$  to  $\ln Z$  for a Yukawa interaction  $\mathcal{L}_I = g \bar{\psi} \psi \phi$ , with a bosonic scalar field  $\phi$  and a fermionic field  $\psi$ . Give  $\langle S_I^2 \rangle_0$  in terms of diagrams as well as in terms of momentum sums. (Note that odd powers in the interaction term such as  $\langle S_I \rangle_0$  vanish. Therefore  $\langle S_I^2 \rangle_0$  is the lowest-order correction.)

[End of 10th lecture, Dec 15th, 2008]

## IX. BOSE-EINSTEIN CONDENSATION OF AN INTERACTING BOSE GAS

### A. Spontaneous symmetry breaking and the Goldstone theorem

In Sec. V we have discussed Bose-Einstein condensation of a non-interacting field. Now, having discussed the basics of an interacting theory, we revisit this effect. The inclusion of interactions is not only more realistic for possible applications, but also will give us a better conceptual understanding of Bose-Einstein condensation. In particular, we shall see that Bose-Einstein condensation is an example of spontaneous symmetry breaking and the Goldstone theorem which are extremely important concepts in various fields of theoretical physics. Another consequence of including interactions is that we can compute the condensate for fixed chemical potential, and not only for fixed charge density, as we have done in Sec. V.

We use the same Lagrangian as in Sec. V, see for instance Eq. (87), i.e., we include a chemical potential, but now also include an interaction term as in the previous section,

$$\mathcal{L} = |(\partial_0 - i\mu)\varphi|^2 - |\nabla\varphi|^2 - m^2|\varphi|^2 - \lambda|\varphi|^4. \quad (281)$$

This Lagrangian is invariant under  $U(1)$  rotations  $\varphi \rightarrow e^{-i\alpha}\varphi$ . As in Sec. V we introduce real fields via

$$\varphi = \frac{1}{\sqrt{2}}(\varphi_1 + i\varphi_2), \quad (282)$$

which leads to the Lagrangian

$$\mathcal{L} = \frac{1}{2} \left[ (\partial_0\varphi_1)^2 + (\partial_0\varphi_2)^2 - (\nabla\varphi_1)^2 - (\nabla\varphi_2)^2 + 2\mu(\varphi_2\partial_0\varphi_1 - \varphi_1\partial_0\varphi_2) + (\mu^2 - m^2)(\varphi_1^2 + \varphi_2^2) - \frac{\lambda}{2}(\varphi_1^2 + \varphi_2^2)^2 \right]. \quad (283)$$

As discussed above for the meson case, we separate the zero-mode  $\phi_i$ , allowing for Bose condensation,  $\varphi_i \rightarrow \varphi_i + \phi_i$  (remember Eq. (88)). Then the Lagrangian becomes

$$\mathcal{L} = -U(|\phi|^2) + \mathcal{L}^{(2)} + \mathcal{L}^{(3)} + \mathcal{L}^{(4)}, \quad (284)$$

with

$$U(|\phi|^2) = \frac{m^2 - \mu^2}{2}(\phi_1^2 + \phi_2^2) + \frac{\lambda}{4}(\phi_1^2 + \phi_2^2)^2, \quad (285a)$$

$$\begin{aligned} \mathcal{L}^{(2)} = & -\frac{1}{2} [ -(\partial_0\varphi_1)^2 - (\partial_0\varphi_2)^2 + (\nabla\varphi_1)^2 + (\nabla\varphi_2)^2 - 2\mu(\varphi_2\partial_0\varphi_1 - \varphi_1\partial_0\varphi_2) \\ & + (m^2 - \mu^2)(\varphi_1^2 + \varphi_2^2) + \lambda(3\phi_1^2 + \phi_2^2)\varphi_1^2 + \lambda(\phi_1^2 + 3\phi_2^2)\varphi_2^2 + 4\lambda\phi_1\phi_2\varphi_1\varphi_2 ], \end{aligned} \quad (285b)$$

$$\mathcal{L}^{(3)} = -\lambda(\phi_1\varphi_1 + \phi_2\varphi_2)(\varphi_1^2 + \varphi_2^2), \quad (285c)$$

$$\mathcal{L}^{(4)} = -\frac{\lambda}{4}(\varphi_1^2 + \varphi_2^2)^2. \quad (285d)$$

As explained in Sec. V, the linear terms in the fields  $\varphi_i$  vanish after space-time integration; hence we have already omitted them here. Remarkably, besides the obvious quartic interaction term  $\mathcal{L}^{(4)}$ , there is an interaction term *cubic* in the fields, induced by the condensate  $\phi_i$ . (Note that here we use the term “condensate” for what is, in the particle physics context, also called “vacuum expectation value” for the field, or shortly “vev”.) Both interaction terms will lead to loop corrections as discussed in the previous section. However, let us first discuss the “tree-level” contributions.

Again, for symmetry reasons, we can choose  $\phi_2 = 0$  and denote  $\phi \equiv \phi_1$ . Then, the potential becomes

$$U(\phi^2) = \frac{m^2 - \mu^2}{2}\phi^2 + \frac{\lambda}{4}\phi^4, \quad (286)$$

and, following the same steps as in Sec. V, the tree-level propagator is

$$D_0^{-1}(K) = \begin{pmatrix} -K^2 + m^2 + 3\lambda\phi^2 - \mu^2 & -2ik_0\mu \\ 2ik_0\mu & -K^2 + m^2 + \lambda\phi^2 - \mu^2 \end{pmatrix}. \quad (287)$$

Note that  $D_0$  knows about the interaction, although we have only considered the terms quadratic in the fields. This is because of the condensate which appears together with the coupling constant  $\lambda$ . Again following the steps in Sec. V we obtain the tree-level thermodynamic potential,

$$\frac{\Omega}{V} = U(\phi^2) + \frac{1}{2V} \text{Tr} \ln \frac{D_0^{-1}(K)}{T^2}. \quad (288)$$

The poles of the tree-level propagator  $D_0$  (i.e., the zeros of the determinant of the inverse propagator  $D_0^{-1}$ ) correspond to the quasiparticle energies, which we denote by  $\epsilon_k^\pm$ . They can be used to compute the  $\text{Tr} \ln$  (which is the same as  $\ln \det$ ). Defining

$$m_1^2 \equiv m^2 + 3\lambda\phi^2, \quad (289a)$$

$$m_2^2 \equiv m^2 + \lambda\phi^2, \quad (289b)$$

we obtain (cf. Eq. (92))

$$\begin{aligned} \ln \det \frac{D_0^{-1}(K)}{T^2} &= \ln \prod_K \frac{1}{T^4} [(-K^2 + m_1^2 - \mu^2)(-K^2 + m_2^2 - \mu^2) - 4\mu^2 k_0^2] \\ &= \ln \prod_K \frac{1}{T^4} [(\epsilon_k^+)^2 - k_0^2][(\epsilon_k^-)^2 - k_0^2] \\ &= \sum_K \left[ \ln \frac{(\epsilon_k^+)^2 - k_0^2}{T^2} + \ln \frac{(\epsilon_k^-)^2 - k_0^2}{T^2} \right]. \end{aligned} \quad (290)$$

The quasiparticle energies are

$$\epsilon_k^\pm = \sqrt{E_k^2 + \mu^2} \mp \sqrt{4\mu^2 E_k^2 + \delta M^4}, \quad (291)$$

where we abbreviated

$$E_k \equiv \sqrt{k^2 + M^2}, \quad (292)$$

and

$$M^2 \equiv \frac{m_1^2 + m_2^2}{2} = m^2 + 2\lambda\phi^2, \quad (293a)$$

$$\delta M^2 \equiv \frac{m_1^2 - m_2^2}{2} = \lambda\phi^2. \quad (293b)$$

Performing the Matsubara sum in Eq. (290) as usual, inserting the result into the thermodynamic potential (288) and dropping the vacuum contribution yields

$$\frac{\Omega}{V} = \frac{m^2 - \mu^2}{2} \phi^2 + \frac{\lambda}{4} \phi^4 + T \sum_{e=\pm} \int \frac{d^3k}{(2\pi)^3} \ln \left( 1 - e^{-\epsilon_k^e/T} \right). \quad (294)$$

The quasiparticle energies (291) look complicated and are best understood through their limit cases. Without interactions,  $\lambda = 0$ , we have  $M = m$  and  $\delta M = 0$ . Consequently, we recover the usual particle and antiparticle dispersions. Also for vanishing chemical potential, the dispersions become very simple,

$$\epsilon_k^\pm = \begin{cases} \sqrt{k^2 + m^2} \mp \mu & \text{for } \lambda = 0 \\ \sqrt{k^2 + m_{2/1}^2} & \text{for } \mu = 0 \end{cases}, \quad (295)$$

with  $m_1^2, m_2^2$  defined in Eqs. (289).

Firstly, let us discuss the zero-temperature case,  $T = 0$ . In this case, the thermodynamic potential is simply the potential  $U$ ,

$$\frac{\Omega(T=0)}{V} = \frac{m^2 - \mu^2}{2} \phi^2 + \frac{\lambda}{4} \phi^4. \quad (296)$$

Minimization of  $\Omega$  with respect to  $\phi$  yields the ground state (corresponding to the state with maximal pressure). For chemical potentials  $|\mu| < m$ , the minimum is at  $\phi = 0$ . In accordance with our observation in the noninteracting theory, this means that there is no condensation in this case. One rather needs a negative coefficient in front of the  $\phi^2$  term for condensation, i.e.,  $|\mu| > m$  (negative “mass parameter squared” in field-theoretical treatments without chemical potential). In this case, the potential has a “mexican hat” or “bottom of a wine bottle” shape (we consider

a repulsive interaction for which  $\lambda > 0$ ). Remember that we started from a complex field  $\varphi$ , hence the rotationally symmetric wine bottle potential. The minimum now is at  $\phi \neq 0$ . Such a minimum cannot be invariant under  $U(1)$ . However, all possible minima are degenerate, i.e., we are free to choose an arbitrary direction (and have already done this by choosing  $\phi = \phi_1$ ). This mechanism, where the Lagrangian has a symmetry which is not respected by the ground state, is called *spontaneous symmetry breaking*, as already briefly mentioned in Sec. V. The object that breaks the symmetry and which is zero in the symmetric phase (here the condensate  $\phi$ ) is called the *order parameter*. Other examples of spontaneous symmetry breaking are: (i) ferromagnetism; in this case, the rotational symmetry  $SO(3)$  is, by alignment of all microscopic spins and hence a nonzero magnetization, broken down to  $U(1)$ . (ii) superconductivity; in this case, the order parameter is a condensate of Cooper pairs, breaking the electromagnetic gauge group.

For  $\mu^2 > m^2$  we can easily solve the minimization condition

$$\frac{\partial\Omega(T=0)}{\partial\phi} = 0 \quad \Rightarrow \quad \phi^2 = \frac{\mu^2 - m^2}{\lambda}. \quad (297)$$

This leads to  $M^2 = 2\mu^2 - m^2$ ,  $\delta M^2 = \mu^2 - m^2$  and thus

$$\epsilon_k^\pm = \sqrt{k^2 + (3\mu^2 - m^2) \mp \sqrt{4\mu^2 k^2 + (3\mu^2 - m^2)^2}}. \quad (298)$$

We see that  $\epsilon_k^+$  becomes gapless, i.e.,  $\epsilon_{k=0}^+ = 0$ . We can expand this mode for small momenta to obtain

$$\epsilon_k^+ \simeq \sqrt{\frac{\mu^2 - m^2}{3\mu^2 - m^2}} k = \sqrt{\frac{M^2 - \mu^2}{M^2 - \mu^2}} k. \quad (299)$$

For  $\mu = 0$ , the slope of this linear dispersion is 1; the chemical potential changes the slope. This gapless mode is called the *Goldstone mode*. Its presence is of great importance since one is very often interested in the low-energy limit of a theory. The Goldstone mode can be excited with arbitrarily small energy, i.e., it will be present no matter how small the considered energy is. Moreover, its presence is a very general fact, due to the *Goldstone theorem* which says that in any system with a spontaneously broken global symmetry there is a gapless mode.<sup>3</sup>

The energy gap of the second mode is given by

$$\epsilon_{k=0}^- = \sqrt{2(3\mu^2 - m^2)} = \sqrt{2(M^2 + \mu^2)}. \quad (300)$$

## B. Symmetry restoration at finite temperature

At a certain temperature, called critical temperature  $T_c$ , the symmetry will be restored, i.e., the ground state will be symmetric under the original symmetry group. In our case this means that for  $T > T_c$  we have  $\phi = 0$  (in the case of a ferromagnet the magnetization vanishes above the critical temperature, there called Curie temperature; in a superconductor, the condensate of Cooper pairs vanishes above  $T_c$  etc.). The general treatment of symmetry restoration and the determination of the critical temperature is complicated. Here we shall first discuss the qualitative picture which gives a good physical understanding for the symmetry restoration process. We shall do so with the help of a very simple approximation. We shall also see that this approximation has obvious problems.

For the sake of simplicity let us set  $\mu = 0$ . This means, in order to have condensation at small temperatures, we need a negative mass parameter squared. Therefore, we introduce the positive square  $c^2$  via

$$m^2 = -c^2. \quad (301)$$

The excitations energies are now much simpler and given in Eq. (295). We shall use the high-temperature approximation  $T^2 \gg c^2$  for the potential in Eq. (294). Then, with

$$\int_0^\infty dx x^2 \ln(1 - e^{-\sqrt{x^2+y^2}}) = -\frac{\pi^4}{45} + \frac{\pi^2}{12} y^2 + \mathcal{O}(y^3), \quad (302)$$

---

<sup>3</sup> Notice the specification *global* symmetry. For a *local* symmetry, for instance in the case of superconductivity, the Goldstone mode is “eaten up” by the gauge fields, giving rise to a magnetic screening mass, the *Meissner mass*.

we can approximate the potential as follows,

$$\begin{aligned}
\frac{\Omega}{V} &= -\frac{c^2}{2}\phi^2 + \frac{\lambda}{4}\phi^4 + T \sum_{\epsilon=\pm} \int \frac{d^3k}{(2\pi)^3} \ln \left( 1 - e^{-\epsilon_k/T} \right) \\
&= -\frac{c^2}{2}\phi^2 + \frac{\lambda}{4}\phi^4 + \frac{T^4}{2\pi^2} \sum_{i=1,2} \int_0^\infty dx x^2 \ln \left( 1 - e^{-\sqrt{x^2+(m_i/T)^2}} \right) \\
&\simeq -\frac{c^2}{2}\phi^2 + \frac{\lambda}{4}\phi^4 - \frac{T^4\pi^2}{45} + \frac{T^2(m_1^2 + m_2^2)}{24}.
\end{aligned} \tag{303}$$

Now, with Eqs. (289) and (301) this becomes, ordered in powers of  $\phi$ ,

$$\frac{\Omega}{V} = \left( -\frac{c^2}{2} + \frac{\lambda T^2}{6} \right) \phi^2 + \frac{\lambda}{4}\phi^4 - \frac{T^4\pi^2}{45} - \frac{c^2 T^2}{12}. \tag{304}$$

This result shows that the coefficient of the quadratic term becomes larger with increasing temperature until it becomes positive for temperatures larger than the *critical temperature*

$$T_c^2 = \frac{3c^2}{\lambda}. \tag{305}$$

This indicates a second-order phase transition to the restored phase. (A first-order phase transition can only occur if there was also a cubic term  $\phi^3$ .) More precisely, the minimum as a function of temperature is

$$\phi^2(T) = \frac{c^2}{\lambda} - \frac{T^2}{3}, \tag{306}$$

for  $T < T_c$  and  $\phi^2 = 0$  for  $T > T_c$ . This is of course a very crude approximation. In particular, we see the following severe problem. With Eqs. (289) and (306) we observe that the excitation energies are given by the following masses (for  $T < T_c$ ),

$$m_1^2 = 2c^2 - \lambda T^2, \tag{307a}$$

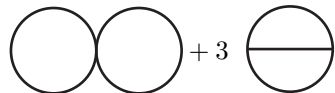
$$m_2^2 = -\frac{\lambda T^2}{3}. \tag{307b}$$

For  $T = 0$  we recover the gapless Goldstone mode and the gapped mode. However, for nonzero temperature, the Goldstone mode acquires an imaginary energy for small momenta which is obviously unphysical. It is no surprise that the applied approximation is incomplete at best, since we know from the previous section that loops can give corrections to the potential of the order of  $\lambda T^2$ . This is exactly the order which was responsible for the symmetry restoration above. Therefore, in the next subsection we improve our result by including loop corrections.

[End of 11th lecture, Jan 12th, 2009]

### C. Including loop corrections

So far we have ignored the contributions from the interaction terms (285c) and (285d) in the Lagrangian. From Sec. VIII A we know that the quartic term gives rise to the “double-bubble” diagram in the thermodynamic potential. The cubic term gives an additional contribution, such that up to two loops we find the contributions (omitting the arrows in the loop diagrams)

$$\ln Z_I^{\text{two loops}} = 3 \left( \text{Diagram 1} \right) + 3 \left( \text{Diagram 2} \right) \tag{308}$$


Note that now each line denotes a  $2 \times 2$  propagator, given in Eq. (287). We shall in the following neglect the second diagram which is justified in the low-momentum, large-temperature approximation that we shall employ below (note that the self-energy from the second diagram depends on momentum, in contrast to the self-energy from the first diagram). For an estimate of the self-energy from the second diagram, solve the following

**Exercise 8:** Show that the momentum-dependent self-energy

$$\Sigma(P) \equiv \phi^2 \lambda^2 \frac{T}{V} \sum_K D_0(P-K) D_0(K) \tag{309}$$

with the propagator  $D_0(K) = 1/(k_0^2 - \epsilon_k^2)$ ,  $\epsilon_k^2 = k^2 + m^2$ , is given in the zero-frequency, low-momentum, and large-temperature limits as

$$\lim_{\mathbf{p} \rightarrow 0} \Sigma(0, \mathbf{p}) \simeq \frac{\phi^2 \lambda^2 T}{2m\pi} \left( \frac{1}{8} + \frac{1}{3\pi} \right). \quad (310)$$

*Hint: Use the result from Exercise 1 for the Matsubara sum.*

To find the result for the first diagram we write the interaction Lagrangian (285d) as

$$\mathcal{L}^{(4)} = -\Lambda_{abcd} \varphi_a \varphi_b \varphi_c \varphi_d, \quad a, b, c, d = 1, 2, \quad (311)$$

with a symmetrized tensor,

$$\Lambda_{abcd} = \frac{\lambda}{12} (\delta_{ab} \delta_{cd} + \delta_{ac} \delta_{bd} + \delta_{ad} \delta_{bc}). \quad (312)$$

The tensor structure comes from the two components of the complex field  $\varphi$  and replaces the scalar  $\lambda$  for the case of a single scalar field. Consequently,

$$\ln Z_I^{(1)} = -3\Lambda_{abcd} \frac{T}{V} \sum_K D_0^{ab}(K) \sum_Q D_0^{cd}(Q), \quad (313)$$

where  $D_0^{ab}$  are the components of the  $2 \times 2$  propagator

$$D_0(K) = \frac{1}{[(\epsilon_k^+)^2 - k_0^2][(\epsilon_k^-)^2 - k_0^2]} \begin{pmatrix} -K^2 + m_2^2 - \mu^2 & 2ik_0\mu \\ -2ik_0\mu & -K^2 + m_1^2 - \mu^2 \end{pmatrix}. \quad (314)$$

Inserting the tensor (312) into Eq. (313) and performing the sum over  $a, b, c, d$  yields

$$\ln Z_I^{(1)} = -\frac{\lambda T}{4V} \left\{ 3 \left[ \sum_Q D_0^{11}(Q) \right]^2 + 3 \left[ \sum_Q D_0^{22}(Q) \right]^2 + 2 \sum_Q D_0^{11}(Q) \sum_K D_0^{22}(K) \right\}, \quad (315)$$

where we have used that the Matsubara sum over the off-diagonal components  $D_0^{12} = -D_0^{21}$  vanishes. Therefore we are left with the diagonal components. The self-energy (which, due to the definition (232) now is also a  $2 \times 2$  matrix) can now be determined from the relation (249). With Eq. (313) we find

$$\begin{aligned} \Pi^{ab} &= -2 \frac{\delta \ln Z_I^{(1)}}{\delta D_0^{ab}} \\ &= 12\Lambda_{abcd} \frac{T}{V} \sum_Q D_0^{cd}(Q) \\ &= \lambda (\delta_{ab} \delta_{cd} + \delta_{ac} \delta_{bd} + \delta_{ad} \delta_{bc}) \frac{T}{V} \sum_Q D_0^{cd}(Q). \end{aligned} \quad (316)$$

Written as a matrix, this is

$$\Pi = \lambda \frac{T}{V} \sum_Q \begin{pmatrix} 3D_0^{11}(Q) + D_0^{22}(Q) & 0 \\ 0 & D_0^{11}(Q) + 3D_0^{22}(Q) \end{pmatrix}. \quad (317)$$

Using two different line styles for the two modes 1 and 2 we can bring Eqs. (315) and (317) into a diagrammatic form,

$$4 \ln Z_I^{(1)} = 3 \left( \text{two solid circles} \right) + 3 \left( \text{two dashed circles} \right) + 2 \left( \text{one solid and one dashed circle} \right), \quad (318)$$



and

$$\Pi^{11} = -3 \frac{\text{---} \bigcirc \text{---}}{\text{---}} - \frac{\text{---} \bigcirc \text{---}}{\text{---}}, \quad (319a)$$

$$\Pi^{22} = -3 \frac{\text{---} \bigcirc \text{---}}{\text{---}} - \frac{\text{---} \bigcirc \text{---}}{\text{---}}. \quad (319b)$$

As a check, we see that these self-energy diagrams arise from cutting the respective lines in Eq. (318). To evaluate the self-energy from Eq. (317) we again set  $\mu = 0$  for simplicity. The result will give us the loop corrections to the masses given in Eq. (307). In the high-temperature approximation,  $T \gg m_1, m_2$ , the propagators of the two modes become identical,  $D_0^{11} \simeq D_0^{22} \simeq -1/K^2$ . Then we can use the results of Sec. VIII C to obtain, after subtracting the vacuum contribution,

$$\lambda \frac{T}{V} \sum_Q D_0^{11}(Q) = \lambda \frac{T}{V} \sum_Q D_0^{22}(Q) = \frac{\lambda T^2}{12}, \quad (320)$$

i.e.,  $\Pi^{11} \simeq \Pi^{22} \simeq \lambda T^2/3$ . Consequently, the corrected masses of the two modes are, for  $T < T_c$ ,

$$m_1^2 + \Pi^{11} = 2 \left( c^2 - \frac{\lambda T^2}{3} \right) = \frac{2\lambda T_c^2}{3} \left( 1 - \frac{T^2}{T_c^2} \right), \quad (321a)$$

$$m_2^2 + \Pi^{22} = 0, \quad (321b)$$

where we have used Eqs. (305) and (307). We see that the Goldstone mode now remains gapless up to  $T = T_c$  as it should be. Moreover, the other mode becomes ungapped at the phase transition. Then, for  $T > T_c$ , both masses become identical,  $m_1^2 = m_2^2 = -c^2 + \lambda T^2/3 \geq 0$ .

The case of a nonvanishing chemical potential is similar. However, in this case, the gapped mode has an energy gap even at  $T = T_c$ , where the energy is  $\mu$  (as opposed to 0 in the case without chemical potential). For a self-consistent treatment of this case see for instance Sec. III in Ref. [5].

## X. THE PHOTON PROPAGATOR IN A QED PLASMA

### A. Photon polarization tensor

From Exercise 8 we know the structure of the self-energy diagrams for fermions interacting via a bosonic field through the Yukawa interaction. In QED, discussed in Sec. VII, electrons interact in a similar way with photons, namely through an interaction term

$$\mathcal{L}_I = e \bar{\psi} \gamma^\mu A_\mu \psi, \quad (322)$$

cf. Eq. (166). From this expression we can construct the one-loop photon self-energy

$$\Pi_{\mu\nu}(Q) = \text{---} \bigcirc \text{---} = e^2 \frac{T}{V} \sum_K \text{Tr}[\gamma_\mu G_0(K) \gamma_\nu G_0(P)], \quad (323)$$

where the trace is taken over Dirac space, where we have abbreviated

$$P \equiv K - Q, \quad (324)$$

and where the electron propagator is given by Eq. (139). A gauge interaction is of course different from a simple Yukawa interaction. We shall come to the subtleties related to gauge invariance in the next subsection. First we

compute the photon self-energy, also called *polarization tensor*. For simplicity, we shall consider the ultrarelativistic limit  $m = 0$  and set the chemical potential to zero,  $\mu = 0$ . Then, the fermion propagator simply becomes

$$G_0(K) = -\frac{\gamma^\mu K_\mu}{K^2}. \quad (325)$$

With the trace

$$\text{Tr}[\gamma_\mu \gamma_\sigma \gamma_\nu \gamma_\rho] = 4(g_{\mu\sigma} g_{\nu\rho} + g_{\mu\rho} g_{\sigma\nu} - g_{\mu\nu} g_{\sigma\rho}), \quad (326)$$

where  $g_{\mu\nu} = \text{diag}(1, -1, -1, -1)$  is the metric tensor, we obtain

$$\text{Tr}[\gamma_\mu \gamma^\sigma K_\sigma \gamma_\nu \gamma^\rho P_\rho] = 4(K_\mu P_\nu + K_\nu P_\mu - g_{\mu\nu} K \cdot P). \quad (327)$$

In the following we need to treat the following three cases separately,

$$\Pi_{00}(Q) = 4e^2 \frac{T}{V} \sum_K \frac{k_0 p_0 + \mathbf{k} \cdot \mathbf{p}}{(k_0^2 - k^2)(p_0^2 - p^2)}, \quad (328a)$$

$$\Pi_{0i}(Q) = \Pi_{i0}(Q) = 4e^2 \frac{T}{V} \sum_K \frac{k_0 p_i + k_i p_0}{(k_0^2 - k^2)(p_0^2 - p^2)}, \quad (328b)$$

$$\Pi_{ij}(Q) = 4e^2 \frac{T}{V} \sum_K \frac{k_i p_j + k_j p_i + \delta_{ij}(k_0 p_0 - \mathbf{k} \cdot \mathbf{p})}{(k_0^2 - k^2)(p_0^2 - p^2)}. \quad (328c)$$

The Matsubara sum over  $k_0$  is taken over fermionic Matsubara frequencies since the loop is an electron loop. Since the external momentum  $Q$  belongs to the photon, the Matsubara frequencies in  $q_0$  are bosonic. Therefore we can use the result from Exercise 4. With the relations

$$1 - f_F(\epsilon_1) - f_F(\epsilon_2) = \frac{f_F(\epsilon_1) f_F(\epsilon_2)}{f_B(\epsilon_1 + \epsilon_2)} = -\frac{f_F(-\epsilon_1) f_F(-\epsilon_2)}{f_B(-\epsilon_1 - \epsilon_2)}, \quad (329a)$$

$$f_F(\epsilon_1) - f_F(\epsilon_2) = \frac{f_F(-\epsilon_1) f_F(\epsilon_2)}{f_B(-\epsilon_1 + \epsilon_2)} = -\frac{f_F(\epsilon_1) f_F(-\epsilon_2)}{f_B(\epsilon_1 - \epsilon_2)}, \quad (329b)$$

we can write Eq. (152) as (changing  $q_0$  to  $-q_0$ )

$$T \sum_{k_0} \frac{(k_0 + \xi_1)(k_0 - q_0 + \xi_2)}{(k_0^2 - \epsilon_1^2)[(k_0 - q_0)^2 - \epsilon_2^2]} = \frac{1}{4\epsilon_1 \epsilon_2} \left\{ \left[ \frac{(\epsilon_1 - \xi_1)(\epsilon_2 - \xi_2)}{q_0 + \epsilon_1 - \epsilon_2} - \frac{(\epsilon_1 + \xi_1)(\epsilon_2 + \xi_2)}{q_0 - \epsilon_1 + \epsilon_2} \right] [f(\epsilon_1) - f(\epsilon_2)] \right. \\ \left. + \left[ \frac{(\epsilon_1 + \xi_1)(\epsilon_2 - \xi_2)}{q_0 - \epsilon_1 - \epsilon_2} - \frac{(\epsilon_1 - \xi_1)(\epsilon_2 + \xi_2)}{q_0 + \epsilon_1 + \epsilon_2} \right] [1 - f(\epsilon_1) - f(\epsilon_2)] \right\}. \quad (330)$$

Since eventually only fermionic distribution functions occur we have abbreviated  $f \equiv f_F$ . Consequently, Eqs. (328) become

$$\Pi_{00} = e^2 \int \frac{d^3 k}{(2\pi)^3} \left[ \left( \frac{1}{q_0 + k - p} - \frac{1}{q_0 - k + p} \right) (1 + \hat{\mathbf{k}} \cdot \hat{\mathbf{p}})(f_k - f_p) \right. \\ \left. + \left( \frac{1}{q_0 - k - p} - \frac{1}{q_0 + k + p} \right) (1 - \hat{\mathbf{k}} \cdot \hat{\mathbf{p}})(1 - f_k - f_p) \right], \quad (331a)$$

$$\Pi_{0i} = -e^2 \int \frac{d^3 k}{(2\pi)^3} \left[ \left( \frac{1}{q_0 + k - p} + \frac{1}{q_0 - k + p} \right) (\hat{p}_i + \hat{k}_i)(f_k - f_p) \right. \\ \left. + \left( \frac{1}{q_0 - k - p} + \frac{1}{q_0 + k + p} \right) (\hat{p}_i - \hat{k}_i)(1 - f_k - f_p) \right], \quad (331b)$$

$$\Pi_{ij} = e^2 \int \frac{d^3 k}{(2\pi)^3} \left[ \left( \frac{1}{q_0 + k - p} - \frac{1}{q_0 - k + p} \right) [\delta_{ij}(1 - \hat{\mathbf{k}} \cdot \hat{\mathbf{p}}) + \hat{k}_i \hat{p}_j + \hat{k}_j \hat{p}_i](f_k - f_p) \right. \\ \left. + \left( \frac{1}{q_0 - k - p} - \frac{1}{q_0 + k + p} \right) [\delta_{ij}(1 + \hat{\mathbf{k}} \cdot \hat{\mathbf{p}}) - \hat{k}_i \hat{p}_j - \hat{k}_j \hat{p}_i](1 - f_k - f_p) \right]. \quad (331c)$$

where we have abbreviated  $f_k \equiv f(k)$  and the unit vector  $\hat{\mathbf{k}} \equiv \mathbf{k}/k$ . We now apply the so-called ‘‘Hard Thermal Loop (HTL)’’ approximation [6], where the dominant contribution comes from fermion momenta  $k \sim T$  (called ‘‘hard’’), and

the photon momentum is of the order of  $q_0$ ,  $q \sim eT$  (called “soft”). In this spirit, we approximate the denominators in Eqs. (331) as (remember  $p = |\mathbf{k} - \mathbf{q}|$ )

$$q_0 \pm k \mp p \simeq q_0 \pm \mathbf{q} \cdot \hat{\mathbf{k}}, \quad q_0 \pm k \pm p \simeq \pm 2k, \quad (332)$$

and the distribution functions as

$$f_k - f_p \simeq \mathbf{q} \cdot \hat{\mathbf{k}} \frac{\partial f_k}{\partial k}, \quad 1 - f_k - f_p \simeq 1 - 2f_k. \quad (333)$$

Now, to compute  $\Pi_{00}$  first note that  $\hat{\mathbf{k}} \cdot \hat{\mathbf{p}} = 1 + \mathcal{O}(q^2/k^2)$ , hence we can neglect the second term. Thus,

$$\begin{aligned} \Pi_{00} &\simeq 2e^2 \int \frac{d^3k}{(2\pi)^3} \left( \frac{1}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}} - \frac{1}{q_0 - \mathbf{q} \cdot \hat{\mathbf{k}}} \right) \mathbf{q} \cdot \hat{\mathbf{k}} \frac{\partial f_k}{\partial k} \\ &= 4e^2 \int \frac{d^3k}{(2\pi)^3} \frac{\mathbf{q} \cdot \hat{\mathbf{k}}}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}} \frac{\partial f_k}{\partial k} \\ &= 4e^2 \int \frac{d^3k}{(2\pi)^3} \left( 1 - \frac{q_0}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}} \right) \frac{\partial f_k}{\partial k}. \end{aligned} \quad (334)$$

In the second step we have used that the angular integral gives the same result for both terms. In the third step we have separated the  $q_0 = 0$  part.<sup>4</sup>

To compute  $\Pi_{i0}$  in the HTL approximation, we can again neglect the second term since  $\hat{p}_i - \hat{k}_i$  is of the order of  $q/k$ , the remaining term, being of order one, is

$$\begin{aligned} \Pi_{i0} &\simeq -2e^2 \int \frac{d^3k}{(2\pi)^3} \left( \frac{1}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}} + \frac{1}{q_0 - \mathbf{q} \cdot \hat{\mathbf{k}}} \right) \mathbf{q} \cdot \hat{\mathbf{k}} \hat{k}_i \frac{\partial f_k}{\partial k} \\ &= 4e^2 \int \frac{d^3k}{(2\pi)^3} \frac{q_0}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}} \hat{k}_i \frac{\partial f_k}{\partial k}. \end{aligned} \quad (335)$$

In  $\Pi_{ij}$ , the second term cannot be neglected. The first term is analogous to the first term in  $\Pi_{00}$  and we obtain

$$\Pi_{ij} = 2e^2 \int \frac{d^3k}{(2\pi)^3} \left[ 2 \left( 1 - \frac{q_0}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}} \right) \hat{k}_i \hat{k}_j \frac{\partial f_k}{\partial k} - \frac{1}{k} (\delta_{ij} - \hat{k}_i \hat{k}_j) (1 - 2f_k) \right]. \quad (336)$$

The 1 in the factor  $1 - 2f_k$  is the vacuum contribution which has to be subtracted as discussed for the case of the self-energy in  $\phi^4$  theory. Then, with

$$\int \frac{d\Omega}{4\pi} \hat{k}_i \hat{k}_j = \frac{\delta_{ij}}{3}, \quad (337)$$

we see via partial integration that only one term in Eq. (336) survives,

$$\int \frac{d^3k}{(2\pi)^3} \left[ \hat{k}_i \hat{k}_j \frac{\partial f_k}{\partial k} + \frac{1}{k} (\delta_{ij} - \hat{k}_i \hat{k}_j) f_k \right] = \frac{\delta_{ij}}{6\pi^2} \int_0^\infty dk \left( k^2 \frac{\partial f_k}{\partial k} + 2k f_k \right) = 0. \quad (338)$$

Consequently, we are left with

$$\Pi_{ij} = -4e^2 \int \frac{d^3k}{(2\pi)^3} \frac{q_0}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}} \hat{k}_i \hat{k}_j \frac{\partial f_k}{\partial k}. \quad (339)$$

In all three results (334), (335), and (339) we have the same  $k$  integral which we can perform exactly,

$$\int_0^\infty dk k^2 \frac{\partial f_k}{\partial k} = -\frac{T^2}{2} \int_0^\infty dx \frac{x^2}{1 + \cosh x} = -\frac{T^2 \pi^2}{6}. \quad (340)$$

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<sup>4</sup> Note that the limits  $q_0 \rightarrow 0$ ,  $q \rightarrow 0$  do not commute. We shall see in the subsequent sections that different limits correspond to different physics.

Consequently, we have

$$\Pi_{00} = -2m^2 \left( 1 - \int \frac{d\Omega}{4\pi} \frac{q_0}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}} \right), \quad (341a)$$

$$\Pi_{0i} = -2m^2 \int \frac{d\Omega}{4\pi} \frac{q_0 \hat{k}_i}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}}, \quad (341b)$$

$$\Pi_{ij} = 2m^2 \int \frac{d\Omega}{4\pi} \frac{q_0 \hat{k}_i \hat{k}_j}{q_0 + \mathbf{q} \cdot \hat{\mathbf{k}}}, \quad (341c)$$

where we defined

$$m^2 \equiv \frac{e^2 T^2}{6}. \quad (342)$$

## B. Photon propagator

After having computed the polarization tensor, we want to use the result to determine the photon modes in the plasma. To this end, we need the photon propagator  $D(Q)$ . In a covariant gauge with gauge-fixing parameter  $\rho$  the free inverse propagator is

$$D_{0,\mu\nu}^{-1}(Q) = Q^2 g_{\mu\nu} - Q_\mu Q_\nu + \frac{Q_\mu Q_\nu}{\rho}. \quad (343)$$

(Cf. the inverse propagator in axial gauge in Eq. (195).) Physical quantities must of course be independent of  $\rho$ . Inversion gives

$$D_{0,\mu\nu}(Q) = \frac{Q^2 g_{\mu\nu} - Q_\mu Q_\nu}{Q^4} + \rho \frac{Q_\mu Q_\nu}{Q^4}. \quad (344)$$

(One can easily check that  $D_0^{\mu\nu} D_{0,\nu\sigma}^{-1} = \delta_{\mu\sigma}$ .) Let us now introduce projection operators  $P_L, P_T$  via

$$P_T^{00} = P_T^{0i} = P_T^{i0} = 0, \quad (345a)$$

$$P_T^{ij} = \delta^{ij} - \hat{q}^i \hat{q}^j, \quad (345b)$$

and

$$P_L^{\mu\nu} = \frac{Q^\mu Q^\nu}{Q^2} - g^{\mu\nu} - P_T^{\mu\nu}. \quad (346)$$

Both  $P_T$  and  $P_L$  are 4-transverse to  $Q$ , i.e.,  $Q_\mu P_L^{\mu\nu} = Q_\mu P_T^{\mu\nu} = 0$ . The projector  $P_T$  is 3-transverse, while  $P_L$  is 3-longitudinal. We have  $P_L^2 = P_L, P_R^2 = P_R, P_L P_R = P_R P_L = 0$ . In terms of these projectors, the photon self-energy can be written as

$$\Pi_{\mu\nu}(Q) = F(Q) P_{L,\mu\nu} + G(Q) P_{T,\mu\nu}, \quad (347)$$

with scalar functions  $F$  and  $G$ . This follows from rotational invariance and the transversality property of the self-energy  $Q_\mu \Pi^{\mu\nu}$  (in non-abelian gauge theories, the structure of the self-energy is more complicated). With  $D^{-1} = D_0^{-1} + \Pi$  we then find

$$D_{\mu\nu}(Q) = \frac{1}{F(Q) - Q^2} P_{L,\mu\nu} + \frac{1}{G(Q) - Q^2} P_{T,\mu\nu} + \rho \frac{Q_\mu Q_\nu}{Q^4}. \quad (348)$$

From our above results, we can now compute  $F$  and  $G$ . To this end, we choose a frame in which  $\mathbf{q}$  points into the  $z$ -direction. Then we have

$$F(Q) = \frac{Q^2}{q_0 q} \Pi_{0z}(Q), \quad G(Q) = \Pi_{xx}(Q). \quad (349)$$

Then, with Eq. (341b) we find

$$\begin{aligned} F(Q) &= -m^2 \frac{Q^2}{q} \int_{-1}^1 dx \frac{x}{q_0 + qx} \\ &= -2m^2 \frac{Q^2}{q^2} \left( 1 - \frac{q_0}{2q} \ln \frac{q_0 + q}{q_0 - q} \right), \end{aligned} \quad (350)$$

and

$$\begin{aligned} G(Q) &= \frac{m^2}{2\pi} \int_0^{2\pi} d\varphi \int_0^\pi d\theta \sin \theta \frac{q_0 \sin^2 \theta \cos^2 \varphi}{q_0 + q \cos \theta} \\ &= \frac{m^2 q_0}{2} \int_{-1}^1 dx \frac{1 - x^2}{q_0 + qx} \\ &= \frac{m^2 q_0}{q} \left( \frac{q_0}{q} - \frac{Q^2}{2q^2} \ln \frac{q_0 + q}{q_0 - q} \right) \\ &= m^2 - \frac{1}{2} F(Q). \end{aligned} \quad (351)$$

In both functions  $F$  and  $G$  the Legendre function of the second kind

$$Q_0(x) = \frac{1}{2} \ln \frac{x+1}{x-1} \quad (352)$$

appears. This function is defined in the complex plane cut from -1 to 1. Consequently,  $F(Q)$  and  $G(Q)$  are defined in the complex  $q_0$  plane cut from  $-q$  to  $q$ . For timelike  $Q^2 > 0$  ( $q_0$  real),  $F$  and  $G$  are real, while for spacelike  $Q^2 < 0$  they become complex.

### C. Debye screening

In order to discuss the physical meaning of the photon propagator with the functions  $F$  and  $G$ , we first consider a static point charge  $\mathcal{Q}$  in the plasma. The resulting potential in position space then is given by the function  $F$  (see chapter 6.3 in Ref. [2] for more details)

$$V(r) = \mathcal{Q} \int \frac{d^3 q}{(2\pi)^3} \frac{e^{i\mathbf{q}\cdot\mathbf{r}}}{q^2 + F(q_0 = 0, \mathbf{q})}. \quad (353)$$

For  $F = 0$  we recover the usual Coulomb potential  $V(r) = \mathcal{Q}/r$ . The effect of  $F$  is to screen this potential,

$$V(r) = \frac{\mathcal{Q}}{r} e^{-r m_D}, \quad (354)$$

with the so-called *Debye screening mass*

$$m_D^2 \equiv F(q_0 = 0, q \rightarrow 0) = -\Pi_{00}(q_0 = 0, q \rightarrow 0) = \frac{e^2 T^2}{3}. \quad (355)$$

This screening is easy to understand: since we are at finite temperature, there are electrons and positrons in the system. Say  $\mathcal{Q}$  is a negative charge, then positrons will be attracted and screen the charge. They do so on a length scale given by the inverse Debye mass. In other words, looking from far away (from distances much larger than  $m_D^{-1}$ ) one cannot see the charge. One has to come closer (up to distances  $\sim m_D^{-1}$  or closer) to resolve the charge.

There is no magnetic screening mass,  $G(q_0 = 0, q \rightarrow 0) = 0$ . The magnetic screening mass becomes non-vanishing in a superconductor ( $\rightarrow$  *Meissner mass*), where magnetic fields are screened.

### D. Plasma oscillations and Landau damping

Next we discuss the collective excitations in the plasma. (See, besides the textbooks by LeBellac and Kapusta, Ref. [7] for details.) To this end, we consider the *spectral density*, given by

$$\rho_{L,T}(Q) \equiv \frac{1}{\pi} \text{Im} D_{L,T}(Q), \quad (356)$$

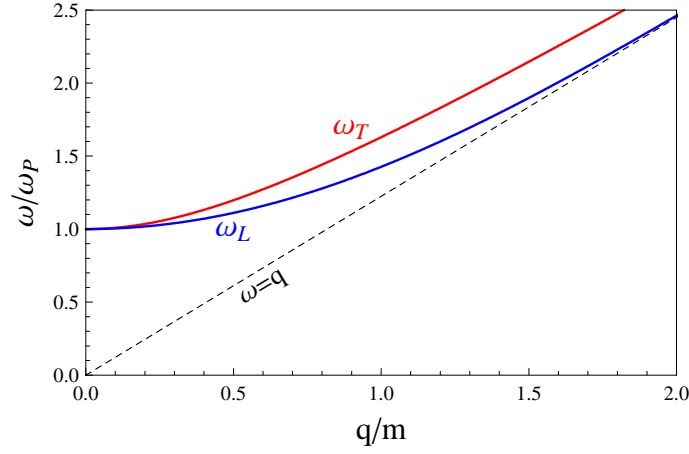


FIG. 1: Longitudinal (plasmon) and transverse modes in a QED plasma.  $\omega_P \equiv eT/3$ ,  $m^2 \equiv e^2 T^2/6$ .

where the longitudinal and transverse parts of the photon propagator are defined as

$$D_L(Q) = \frac{Q^2}{q^2} \frac{1}{F(Q) - Q^2} = \frac{1}{\Pi_{00}(Q) - q^2}, \quad (357a)$$

$$D_T(Q) = \frac{1}{G(Q) - Q^2}, \quad (357b)$$

where we used  $\Pi_{00} = Fq^2/Q^2$  which follows from Eq. (347). The imaginary part in Eq. (356) has to be understood as the imaginary part of the retarded propagator,  $\lim_{\eta \rightarrow 0} \text{Im} D_{L,T}(q_0 + i\eta, \mathbf{q})$ ,  $\eta > 0$ . We then have

$$\rho_L(Q) = \frac{Q^2}{q^2} \frac{1}{\pi} \frac{\text{Im} F}{(\text{Re} F - Q^2)^2 + (\text{Im} F)^2}, \quad (358a)$$

$$\rho_T(Q) = \frac{1}{\pi} \frac{\text{Im} G}{(\text{Re} G - Q^2)^2 + (\text{Im} G)^2}. \quad (358b)$$

As discussed above, the imaginary parts of  $F$  and  $G$  vanish if and only if the four-momentum  $Q$  is spacelike. In this case, the spectral densities vanish except for the case where the whole denominator vanishes. In other words, we use the representation of the delta-function

$$\lim_{\eta \rightarrow 0} \frac{1}{\pi} \frac{\eta}{x^2 + \eta^2} = \delta(x), \quad (359)$$

to obtain

$$\rho_L(Q) = \delta(\text{Re} \Pi_{00} - q^2) + \Theta(q^2 - q_0^2) \frac{1}{\pi} \frac{\text{Im} \Pi_{00}}{(\text{Re} \Pi_{00} - q^2)^2 + (\text{Im} \Pi_{00})^2}, \quad (360a)$$

$$\rho_T(Q) = \delta(\text{Re} G - Q^2) + \Theta(q^2 - q_0^2) \frac{1}{\pi} \frac{\text{Im} G}{(\text{Re} G - Q^2)^2 + (\text{Im} G)^2}. \quad (360b)$$

Since  $\Pi_{00}$  and  $G$  are both even in  $q_0$  we can denote the zeros of the arguments of the delta-functions as  $\pm\omega_L$  and  $\pm\omega_T$ . They correspond to quasiparticles with dispersion  $\omega_{L,T}(q)$ . In general, they have to be determined numerically, see Fig. 1. We see that besides the transverse photon, in a medium there is another, longitudinal, degree of freedom. This quasiparticle is called *plasmon*. Next, we use the general formula

$$\delta[f(x)] = \sum_{x_0} \frac{\delta(x - x_0)}{|f'(x_0)|}, \quad (361)$$

where  $x_0$  are the zeros of the function  $f(x)$ . Moreover, we notice that

$$\frac{\partial \Pi_{00}}{\partial q_0} = \frac{1}{q_0} \left( \Pi_{00} - \frac{3\omega_P^2 q^2}{Q^2} \right), \quad (362)$$

where we have defined the plasma frequency

$$\omega_P^2 = \frac{e^2 T^2}{9}. \quad (363)$$

Thus

$$\left. \frac{\partial \Pi_{00}}{\partial q_0} \right|_{q_0 = \pm \omega_L} = \pm \frac{q^2 (\omega_L^2 - q^2 - 3\omega_P^2)}{\omega_L (\omega_L^2 - q^2)}. \quad (364)$$

And, analogously, for the transverse component,

$$\frac{\partial (G - q_0^2)}{\partial q_0} = \frac{1}{q_0} \left[ \frac{(G - m^2)(3q_0^2 - q^2)}{Q^2} - 2q_0^2 + m^2 \right], \quad (365)$$

and thus

$$\left. \frac{\partial (G - q_0^2)}{\partial q_0} \right|_{q_0 = \pm \omega_T} = \pm \frac{(\omega_T^2 - q^2)^2 - 3\omega_T^2 \omega_P^2}{\omega_T (\omega_T^2 - q^2)}. \quad (366)$$

Consequently, the spectral densities from Eqs. (360) become

$$\rho_L(Q) = \frac{\omega_L (\omega_L^2 - q^2)}{q^2 (3\omega_P^2 + q^2 - \omega_L^2)} [\delta(q_0 - \omega_L) + \delta(q_0 + \omega_L)] + \frac{1}{\pi} \frac{\Theta(q^2 - q_0^2) \text{Im } \Pi_{00}}{(\text{Re } \Pi_{00} - q^2)^2 + (\text{Im } \Pi_{00})^2}, \quad (367a)$$

$$\rho_T(Q) = \frac{\omega_T (\omega_T^2 - q^2)}{3\omega_P^2 \omega_T^2 - (\omega_T^2 - q^2)^2} [\delta(q_0 - \omega_T) + \delta(q_0 + \omega_T)] + \frac{1}{\pi} \frac{\Theta(q^2 - q_0^2) \text{Im } G}{(\text{Re } G - Q^2)^2 + (\text{Im } G)^2}. \quad (367b)$$

We finally remark that the spacelike part of the spectral functions describes *Landau damping*. This is related to scattering processes of the photon off electrons and positrons in the plasma. Through these processes energy is dissipated, i.e., the photon is “damped”.

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