

# Introduction to Electrodynamics by David J. Griffiths Notes

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# 1 Vector Algebra

## 1.6 The Theory of Vector Fields

### 1.6.1 The Helmholtz Theorem

- The **Helmholtz theorem** states that a vector field  $\mathbf{F}$  is uniquely determined if you're given its divergence  $\nabla \cdot \mathbf{F}$ , curl  $\nabla \times \mathbf{F}$ , and sufficient boundary conditions.

### 1.6.2 Potentials

- If the curl of a vector field vanishes everywhere, then it can be expressed as the gradient of a **scalar potential**

$$\nabla \times \mathbf{F} = \mathbf{0} \Leftrightarrow \mathbf{F} = -\nabla V.$$

- If the divergence of a vector field vanishes everywhere, then it can be expressed as the curl of a **vector potential**

$$\nabla \cdot \mathbf{F} = 0 \Leftrightarrow \mathbf{F} = \nabla \times \mathbf{A}.$$

## 2 Electrostatics

### 2.1 The Electric Field

#### 2.1.2 Coulomb's Law

- **Coulomb's law** gives the force between two point charges  $q$  and  $Q$

$$\mathbf{F} = \frac{1}{4\pi\epsilon_0} \frac{qQ}{r^2} \hat{\mathbf{r}}$$

where

$$\epsilon_0 = 8.85 \times 10^{-12} \text{ C}^2/(\text{N m}^2)$$

is the **permittivity of free space** and  $\mathbf{r}$  is the separation vector between the two charges.

#### 2.1.3 The Electric Field

- The **electric field**  $\mathbf{E}$  is a vector field that varies from point to point and gives the force per unit charge that would be exerted on a test charge if placed at a particular point.
- For a collection of  $n$  source charges  $q_i$  at displacements  $\mathbf{r}_i$  from a test charge, the electric field is

$$\mathbf{E} = \frac{1}{4\pi\epsilon_0} \sum_{i=1}^n \frac{q_i}{r_i^2} \hat{\mathbf{r}}_i.$$

#### 2.1.4 Continuous Charge Distributions

- Coulomb's law for a continuous charge distribution is

$$\mathbf{E} = \frac{1}{4\pi\epsilon_0} \int \frac{1}{r^2} \hat{\mathbf{r}} dq.$$

## 2.2 Divergence and Curl of Electrostatic Fields

### 2.2.1 Field Lines, Flux, and Gauss's Law

- **Gauss's law** states that the electric field flux through a closed surface is proportional to the amount of charge within that surface

$$\oint \mathbf{E} \cdot d\mathbf{a} = \frac{1}{\epsilon_0} Q_{\text{enc}}$$

or

$$\nabla \cdot \mathbf{E} = \frac{1}{\epsilon_0} \rho.$$

### 2.2.4 The Curl of $\mathbf{E}$

- The curl of an electric field is  $\mathbf{0}$

$$\nabla \times \mathbf{E} = \mathbf{0}.$$

## 2.3 Electric Potential

### 2.3.1 Introduction to Potential

- The **electric potential** at a point  $\mathbf{r}$  is defined as

$$V(\mathbf{r}) = - \int_{\mathcal{O}}^{\mathbf{r}} \mathbf{E} \cdot d\mathbf{l}$$

where  $\mathcal{O}$  is an agreed origin.

- The potential difference between two points  $\mathbf{a}$  and  $\mathbf{b}$  is

$$V(\mathbf{b}) - V(\mathbf{a}) = - \int_{\mathbf{a}}^{\mathbf{b}} \mathbf{E} \cdot d\mathbf{l}.$$

- The electric field and potential are also related by the equation

$$\mathbf{E} = -\nabla V.$$

### 2.3.2 Comments on Potential

- The choice of origin  $\mathcal{O}$  in the definition of vector potential only affects the absolute potential values, not potential differences. Typically it is chosen to be “at infinity” unless the charge distribution itself extends to infinity.
- Electric potential obeys the superposition principle.
- The units of electric potential is  $\text{N m/C} = \text{J/C} = \text{V}$ .

### 2.3.3 Poisson’s Equation and Laplace’s Equation

- If

$$\nabla \cdot \mathbf{E} = \frac{\rho}{\epsilon_0}$$

and

$$\mathbf{E} = -\nabla V$$

then

$$\begin{aligned} \nabla \cdot (-\nabla V) &= \frac{\rho}{\epsilon_0} \\ \nabla^2 V &= -\frac{\rho}{\epsilon_0}. \end{aligned}$$

This is known as **Poisson’s equation**. In regions where  $\rho = 0$  it reduces to **Laplace’s equation**

$$\nabla^2 V = 0.$$

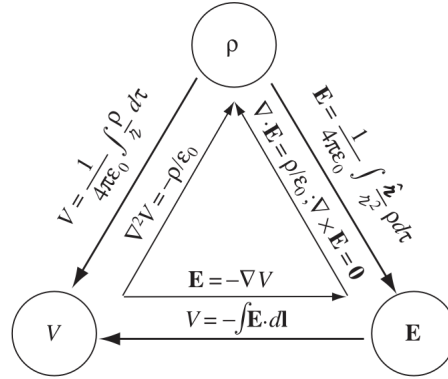
### 2.3.4 The Potential of a Localized Charge Distribution

- The potential of a continuous charge distribution is

$$V(\mathbf{r}) = \frac{1}{4\pi\epsilon_0} \int \frac{\rho(\mathbf{r}')}{r} d\tau'$$

where the reference point is set to infinity.

### 2.3.5 Boundary Conditions



- The normal component of the electric field is discontinuous by an amount  $\sigma/\epsilon_0$  at any boundary, i.e.

$$E_{\text{above}} - E_{\text{below}} = \frac{\sigma}{\epsilon_0}.$$

- The tangential component of the electric field is always continuous at any boundary.
- The electric potential is always continuous at any boundary, however because  $\mathbf{E} = -\nabla V$ , the gradient of the electric potential inherits the discontinuity at boundaries with surface charge, i.e.

$$\nabla V_{\text{above}} - \nabla V_{\text{below}} = -\frac{\sigma}{\epsilon_0} \hat{\mathbf{n}}$$

or

$$\frac{\partial V_{\text{above}}}{\partial n} - \frac{\partial V_{\text{below}}}{\partial n} = -\frac{\sigma}{\epsilon_0}$$

where

$$\frac{\partial V}{\partial n} = \nabla V \cdot \hat{\mathbf{n}}.$$

## 2.4 Work and Energy in Electrostatics

### 2.4.1 The Work It Takes to Move a Charge

- The work required to move a charge  $Q$  from infinity to a point  $\mathbf{r}$  is

$$W = Q[V(\mathbf{r}) - V(\infty)] = QV(\mathbf{r}).$$

In that sense, the electric potential is the energy per unit charge required to assemble a system of point charges.

### 2.4.2 The Energy of a Point Charge Distribution

- If you bring a first charge in from infinity you do no work because there are no other charges. If you bring a second charge in from infinity you do work against the electric field of the first charge. The third does work against the first and second, and so on. Thus the total work required to assemble a collection of charges is

$$\begin{aligned} W &= \frac{1}{4\pi\epsilon_0} \left( \frac{q_1 q_2}{r_{12}} + \frac{q_1 q_3}{r_{13}} + \dots + \frac{q_1 q_n}{r_{1n}} + \frac{q_2 q_3}{r_{23}} + \dots \right) \\ &= \frac{1}{4\pi\epsilon_0} \sum_{i=1}^n \sum_{j>i}^n \frac{q_i q_j}{r_{ij}} \end{aligned}$$

or if we count each pair of charges twice and divide by two

$$W = \frac{1}{8\pi\epsilon_0} \sum_{i=1}^n \sum_{j \neq i}^n \frac{q_i q_j}{r_{ij}}.$$

If we pull  $q_i$  out the front we get

$$W = \frac{1}{2} \sum_{i=1}^n q_i \left( \sum_{j \neq i}^n \frac{1}{4\pi\epsilon_0 r_{ij}} \right) = \frac{1}{2} \sum_{i=1}^n q_i V(\mathbf{r}_i).$$

### 2.4.3 The Energy of a Continuous Charge Distribution

- For a volume charge density  $\rho$  the work to assemble a continuous charge distribution is

$$W = \frac{1}{2} \int \rho V d\tau$$

or equivalently

$$W = \frac{\epsilon_0}{2} \int E^2 d\tau$$

where the integral is taken over all space.

#### 2.4.4 Comments on Electrostatic Energy

- The energy of an electrostatic field does not obey the superposition principle.

### 2.5 Conductors

#### 2.5.1 Basic Properties

- $\mathbf{E} = \mathbf{0}$  inside a conductor because any net electric field causes charges to move, resulting in an induced charge that cancels the electric field.
- $\rho = 0$  inside a conductor. By Gauss's law, if  $\mathbf{E} = \mathbf{0}$  then  $\rho$  must also be 0.
- Any net charge resides on the surface of a conductor.
- A conductor is an equipotential, i.e. the electric potential is the same everywhere in the conductor.
- $\mathbf{E}$  is perpendicular to the surface of the conductor. If there was a tangential component, charge would move to cancel it.

#### 2.5.2 Induced Charges

- If you hold a charge near an uncharged conductor, the two will attract one another. This is because the external charge causes charges within the conductor to move in order to cancel its magnetic field — like charges are repelled and unlike charges are attracted. This results in unlike charges being closer to the external charge and like charges being further away, resulting in a net attraction.
- If you place a charge in a cavity within a conductor, a charge will be induced on the inner surface of the conductor to negate the charge's field within the conductor. If you apply Gauss's law to a Gaussian surface that contains the inner charge and inner surface but not the outer surface, the enclosed charge must be 0. Thus there must be an induced charge of  $q$  on the outer surface which communicates the presence of the inner charge to the outside world.
- If a cavity within a conductor contains no charge,  $\mathbf{E} = \mathbf{0}$  within the cavity.

#### 2.5.3 Surface Charge and Force on a Conductor

- Because the field inside a conductor is zero, the boundary condition  $E_{\text{above}} - E_{\text{below}} = \sigma/\epsilon_0$  requires that the field immediately outside the conductor is

$$\mathbf{E} = \frac{\sigma}{\epsilon_0} \hat{\mathbf{n}}.$$

In terms of potential this is

$$\sigma = -\epsilon_0 \frac{\partial V}{\partial n}.$$

- In the presence of an electric field, a surface charge will experience a force per unit area

$$\mathbf{f} = \sigma \mathbf{E}_{\text{average}} = \frac{1}{2} \sigma (\mathbf{E}_{\text{above}} + \mathbf{E}_{\text{below}}).$$

This applies to any surface charge, however for a conductor  $\mathbf{E}_{\text{below}} = \mathbf{0}$  so

$$\mathbf{f} = \frac{1}{2} \sigma \mathbf{E}_{\text{above}} = \frac{\sigma^2}{2\epsilon_0} \hat{\mathbf{n}}.$$

Notice that this is independent of the sign of  $\sigma$ . This results in an outward **electrostatic pressure** on the surface

$$P = \frac{\epsilon_0}{2} E^2$$

tending to draw the conductor into the field.

#### 2.5.4 Capacitors

- If you have two conductors and place a charge of  $+Q$  on one and  $-Q$  on the other, an electric field will be induced between them. By Coulomb's law the electric field is proportional to the charge so doubling the charge doubles the electric field and thus the electric potential. The constant of proportionality between the charge and the potential difference between the conductors is called their **capacitance**

$$C = \frac{Q}{V}$$

the unit of which is **farads** (F).

- The work required to fully charge a capacitor to potential difference  $V$  is

$$W = \frac{1}{2} CV^2.$$

## 3 Potentials

### 3.1 Laplace's Equation

#### 3.1.1 Introduction

- The primary task of electrostatics is to find the electric field of a stationary charge distribution. In theory this is possible using Coulomb's law, but the integrals are often too difficult. It's often easier to first calculate the potential, but sometimes these too are too difficult. In those cases we can instead try to solve Poisson's equation

$$\nabla^2 V = -\frac{\rho}{\epsilon_0}$$

or Laplace's equation

$$\nabla^2 V = 0.$$



### 3.1.2 Laplace's Equation in One Dimension

- The solution to Laplace's equation in one dimension

$$\frac{d^2V}{dx^2} = 0$$

is

$$V = mx + b.$$

- Under this solution  $V(x)$  is the average of  $V(x + a)$  and  $V(x - a)$  for any  $a$

$$V(x) = \frac{1}{2}[V(x + a) + V(x - a)].$$

- Laplace's equation tolerates no maxima or minima between the endpoints.

### 3.1.3 Laplace's Equation in Two Dimensions

- If you draw a circle of radius  $R$  about a point the average value of  $V$  on the circle is equal to the value at the centre

$$V = \frac{1}{2\pi R} \oint_{\text{circle}} V dl.$$

- $V$  has no maxima or minima — all extrema occur at the boundaries.

### 3.1.4 Laplace's Equation in Three Dimensions

- The value of  $V$  at a point  $\mathbf{r}$  is the average value of  $V$  over a spherical surface of radius  $R$  centred at  $\mathbf{r}$

$$V(\mathbf{r}) = \frac{1}{4\pi R^2} \oint_{\text{sphere}} V da.$$

- $V$  has no maxima or minima — all extrema occur at the boundaries.

### 3.1.5 Boundary Conditions and Uniqueness Theorems

- The **first uniqueness theorem** states that the solution to Laplace's equation in some volume  $\mathcal{V}$  is uniquely determined if  $V$  is specified on the boundary surface  $\mathcal{S}$ .
- A corollary to the first uniqueness theorem is that the potential in a volume  $\mathcal{V}$  is uniquely determined if (a) the charge density throughout the region, and (b) the value of  $V$  on all boundaries, are specified.

### 3.1.6 Conductors and the Second Uniqueness Theorem

- The **second uniqueness theorem** states that in a volume  $\mathcal{V}$  surrounded by conductors and containing a specified charge density  $\rho$ , the electric field is uniquely determined if the total charge on each conductor is given.

## 3.2 The Method of Images

### 3.2.1 The Classic Image Problem

- By the first uniqueness theorem for solutions to Poisson's equation and Laplace's equation, if we can construct a solution that meets the boundary conditions in the region of interest even if it's different outside the region of interest, that's the solution.
- For example, if a point charge  $q$  is placed a distance  $a$  above an infinite grounded conducting plane it will induce some charge in the plane so it's difficult to calculate the potential. However if we consider an alternate scenario where there's a second point charge  $-q$  a distance  $a$  below the plane and the plane is removed, the potential is much easier to calculate. Because the boundary conditions in the region  $z > 0$  are the same, this potential must be a solution to the original problem.

### 3.2.2 Induced Surface Charge

- In the problem above, the surface charge induced on the plane can be calculated as

$$\sigma = -\epsilon_0 \left. \frac{\partial V}{\partial z} \right|_{z=0}.$$

### 3.2.3 Force and Energy

- As potential is the same under the method of images, so is force. Energy however is not the same.

### 3.2.4 Other Image Problems

- In general, **the method of images** can be applied when a stationary charge distribution is in the vicinity of an infinite grounding plane by mirroring the charges across the plane and negating their charges.
- You can't place image charges in the region where you're trying to calculate the potential — that would change  $\rho$  and thus you're solving Poisson's equation with different sources.

## 3.3 Separation of Variables

- The method of separation of variables can be used to solve Laplace's equation when the potential  $V$  or the charge density  $\sigma$  is specified on the boundaries of some region.

### 3.3.2 Spherical Coordinates

- We assume the problems have **azimuthal symmetry** so that  $V$  is independent of  $\phi$ .
- The solutions to the equation

$$\frac{d}{d\theta} \left( \sin \theta \frac{d\Theta}{d\theta} \right) = -l(l+1) \sin \theta \Theta$$

are the **Legendre polynomials** in the variable  $\cos \theta$

$$\Theta(\theta) = P_l(\cos \theta).$$

Although there should be two solutions to  $\Theta(\theta)$  for each value of  $l$ , the second set of solutions “blow up” at  $\theta = 0$  and/or  $\theta = \pi$  so can be excluded on physical grounds. If those values of  $\theta$  are excluded from the region of interest, then these second solutions must be considered.

- Thus, in the case of azimuthal symmetry, the most general separable solution to Laplace’s equations is

$$V(r, \theta) = \sum_{l=0}^{\infty} \left( Ar^l + \frac{B}{r^{l+1}} \right) P_l(\cos \theta).$$

## 3.4 Multipole Expansion

### 3.4.1 Approximate Potentials at Large Distances

- A **physical electric dipole** consists of two equal and opposite charges  $\pm q$  separated by a distance  $d$ .
- The potential of a dipole at distance  $r \gg d$  is

$$V(\mathbf{r}) = \frac{1}{4\pi\epsilon_0} \frac{qd \cos \theta}{r^2}$$

where  $\theta$  is the angle between the line connecting the charges and the line from the centre of the charges to  $\mathbf{r}$ .

- If we place a pair of dipoles next to each other to make a **quadrupole**, the potential goes like  $1/r^3$ . A pair of quadrupoles back to back makes an **octopole**, and the potential goes like  $1/r^4$ , etc.
- The **multipole expansion** of a charge distribution is an expansion of its potential in terms of powers of  $1/r$

$$V(\mathbf{r}) = \frac{1}{4\pi\epsilon_0} \sum_{n=0}^{\infty} \frac{1}{r^{(n+1)}} \int (r')^n P_n(\cos \alpha) \rho(\mathbf{r}') d\tau'.$$

### 3.4.2 The Monopole and Dipole Terms

- The dipole term in the multipole expansion is

$$V_{\text{dip}}(\mathbf{r}) = \frac{1}{4\pi\epsilon_0} \frac{1}{r^2} \int r' \cos \alpha \rho(\mathbf{r}') d\tau'$$

or, since  $\cos \alpha$  is the angle between  $\mathbf{r}$  and  $\mathbf{r}'$

$$V_{\text{dip}}(\mathbf{r}) = \frac{1}{4\pi\epsilon_0} \frac{1}{r^2} \hat{\mathbf{r}} \cdot \int \mathbf{r}' \rho(\mathbf{r}') d\tau'.$$

This integral (which doesn't depend on  $\mathbf{r}$ ) is called the **dipole moment** of the distribution

$$\mathbf{p} = \int \mathbf{r}' \rho(\mathbf{r}') d\tau',$$

and the dipole term simplifies to

$$V_{\text{dip}}(\mathbf{r}) = \frac{1}{4\pi\epsilon_0} \frac{\mathbf{p} \cdot \hat{\mathbf{r}}}{r^2}.$$

- The dipole moment for a collection of point charges is

$$\mathbf{p} = \sum^n q_i \mathbf{r}_i'$$

and for physical dipole is

$$\mathbf{p} = q\mathbf{d}$$

where  $\mathbf{d}$  is the vector from the negative charge to the positive charge.

- A **perfect dipole** is one whose potential is described exactly by the dipole term (no higher terms are needed). This occurs when  $d \rightarrow 0$  and  $q \rightarrow \infty$  while maintaining  $qd = p$ .
- Dipole moments are vectors and add accordingly, e.g. if you have two dipoles  $\mathbf{p}_1$  and  $\mathbf{p}_2$  the total dipole is  $\mathbf{p}_1 + \mathbf{p}_2$ .

### 3.4.3 Origin of Coordinates in Multipole Expansions

- Because the multipole expansion is a power series in inverse powers of  $r$  (the distance to the origin), moving the charge or the origin can affect the expansion. For example, a monopole  $q$  at the origin has potential  $q/4\pi\epsilon_0 r$  — exactly the monopole term in the multipole expansion — but if it is moved a distance  $d$  from the origin it gains a dipole moment  $qd$ .
- If the total charge is 0 the dipole moment doesn't change when you move the distribution or the origin.

### 3.4.4 The Electric Field of a Dipole

- The electric field of a dipole with  $\mathbf{p}$  pointing in the  $z$  direction is

$$\mathbf{E}_{\text{dip}}(r, \theta) = \frac{1}{4\pi\epsilon_0} \frac{p}{r^3} (2 \cos \theta \hat{\mathbf{r}} + \sin \theta \hat{\boldsymbol{\theta}}).$$

## 4 Electric Fields in Matter

### 4.1 Polarization

#### 4.1.1 Dielectrics

- **Conductors** have an effectively unlimited number of free electrons — typically one or two per atom aren't bound and can move around.
- **Insulators** or **dielectrics** have no free electrons — they're all bound to atoms.

#### 4.1.2 Induced Dipoles

- When a neutral atom is placed in an electric field  $\mathbf{E}$  the positive nucleus experiences a force in the direction of  $\mathbf{E}$  and the negative electron cloud experiences a force in the opposite direction to  $\mathbf{E}$ . This causes them to separate slightly and the atom is said to be **polarized**. This also results in a small dipole moment

$$\mathbf{p} = \alpha \mathbf{E}$$

where  $\alpha$  is called the **atomic polarizability**.

- Molecules may have different atomic polarizability constants in different directions. In that case, to calculate the resulting dipole moment you must decompose  $\mathbf{E}$  into the relevant directions and multiply them by the relevant constants.
- The most general equation for the induced dipole moment of a molecule is

$$\begin{aligned} p_x &= \alpha_{xx} E_x + \alpha_{xy} E_y + \alpha_{xz} E_z \\ p_y &= \alpha_{yx} E_x + \alpha_{yy} E_y + \alpha_{yz} E_z \\ p_z &= \alpha_{zx} E_x + \alpha_{zy} E_y + \alpha_{zz} E_z \end{aligned}$$

where the set of nine constants  $\alpha_{ij}$  constitute the **polarizability tensor**. The values of the polarizability tensor depend on the axes you choose, but it's always possible to choose axes such that it is a diagonal matrix.

### 4.1.3 Alignment of Polar Molecules

- Some molecules have a “built-in” dipole moment. These are called **polar molecules**.
- A dipole  $\mathbf{p} = q\mathbf{d}$  in a uniform electric field  $\mathbf{E}$  experiences a torque

$$\boldsymbol{\tau} = \mathbf{p} \times \mathbf{E}.$$

This aligns  $\mathbf{p}$  with  $\mathbf{E}$ .

- A dipole  $\mathbf{p}$  in a nonuniform electric field  $\mathbf{E}$  also experiences a net force

$$\mathbf{F} = \mathbf{F}_+ + \mathbf{F}_- = q(\mathbf{E}_+ - \mathbf{E}_-) = q(\Delta\mathbf{E})$$

where  $\Delta\mathbf{E}$  is the difference in electric field between the ends of the dipole. If the dipole is very short then

$$\mathbf{F} = (\mathbf{p} \cdot \nabla)\mathbf{E}.$$

- The energy of an ideal dipole  $\mathbf{p}$  in an electric field  $\mathbf{E}$  is

$$U = -\mathbf{p} \cdot \mathbf{E}.$$

### 4.1.4 Polarization

- When an external electric field is applied to a material that consists of neutral atoms or nonpolar molecules, the field will induce in each a small dipole moment pointing in the same direction as the field. If the material consists of polar molecules each will experience a torque aligning it with the field. The net result is that there are many small dipoles pointing in the same direction — the material becomes **polarized**. A measure of this is **polarization**  $\mathbf{P}$  which is the dipole moment per unit volume.