



Anisotropic Elasticity

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27-750

Texture, Microstructure & Anisotropy

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Last revised: 7th Feb. '14

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Notation

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F Stimulus (field)
 R Response
 P Property
 \mathbf{j} electric current
 E electric field
 D electric polarization
 ε Strain
 σ Stress (or conductivity)
 ρ Resistivity
 d piezoelectric tensor
 C elastic stiffness
 S elastic compliance

a transformation matrix
 W work done (energy)
 dW work increment
 I identity matrix
 O symmetry operator (matrix)
 Y Young's modulus
 δ Kronecker delta
 e axis (unit) vector
 T tensor
 α direction cosine

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- The objective of this lecture is to provide a mathematical framework for the description of properties, especially when they vary with direction.
- A basic property that occurs in almost all applications is *elasticity*. Although elastic response is *linear* for all practical purposes, it is often *anisotropic* (composites, textured polycrystals etc.).
- Why do we care about elastic anisotropy? In composites, especially fibre composites, it is easy to design in substantial anisotropy by varying the lay-up of the fibres. See, for example: http://www.jwave.vt.edu/crcd/kriz/lectures/Geom_3.html
- Geologists are very familiar with elastic anisotropy and exploit it for understanding seismic results.

In Class Questions

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1. Why is plastic yielding a non-linear property, in contrast to elastic deformation?
2. What is the definition of a tensor?
3. Why is stress is 2nd-rank tensor?
4. Why is elastic stiffness a 4th-rank tensor?
5. What is “matrix notation” (in the context of elasticity)?
6. What are the relationships between tensor and matrix coefficients for stress? Strain? Stiffness? Compliance?
7. Why do we need factors of 2 and 4 in some of these conversion factors?
8. How do we use crystal symmetry to decrease the number of coefficients needed to describe stiffness and compliance?
9. How many independent coefficients are needed for stiffness (and compliance) in cubic crystals? In isotropic materials?
10. How do we express the directional dependence of Young’s modulus?
11. What is Zener’s anisotropy factor?

Anisotropy: Practical Applications

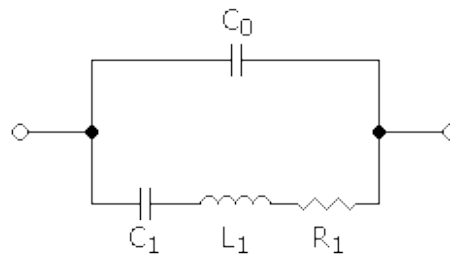
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- The practical applications of anisotropy of composites, especially fiber-reinforced composites are numerous.
- The stiffness of fiber composites varies tremendously with direction. Torsional rigidity is very important in car bodies, boats, aeroplanes etc.
- Even in monolithic polymers (e.g. drawn polyethylene) there exists large anisotropy because of the alignment of the long-chain molecules.

Application example: quartz oscillators

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- Piezoelectric quartz crystals are commonly used for frequency control in watches and clocks. Despite having small values of the piezoelectric coefficients, quartz has positive aspects of low losses and the availability of orientations with negligible temperature sensitivity. The property of **piezoelectricity** relates **strain** to **electric field**, or polarization to stress.
- $\epsilon_{ij} = d_{ijk} E_k$
- PZT, lead zirconium titanate $\text{PbZr}_{1-x}\text{Ti}_x\text{O}_3$, is another commonly used piezoelectric material.



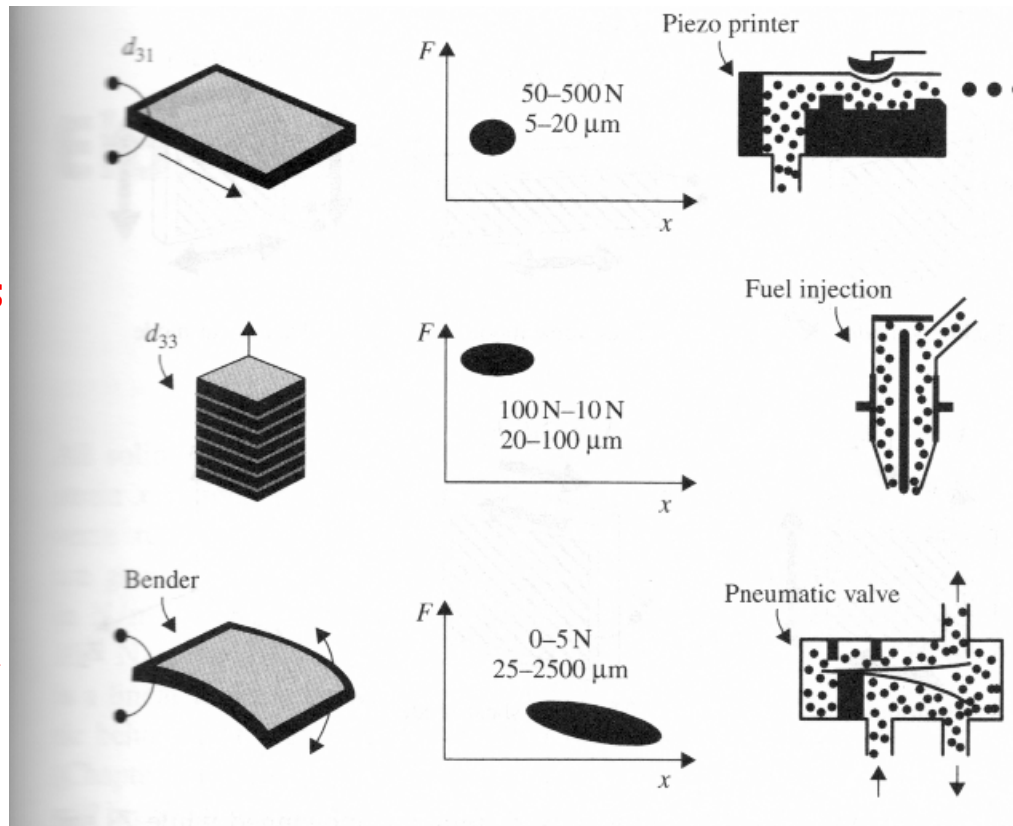
Piezoelectric Devices

- The property of **piezoelectricity** relates **strain** to **electric field**, or polarization to stress.

$$\varepsilon_{ij} = d_{ijk} E_k$$

- PZT, lead zirconium titanate $\text{PbZr}_{1-x}\text{Ti}_x\text{O}_3$, is another commonly used piezoelectric material.

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Note: Newnham consistently uses vector-matrix notation, rather than tensor notation. We will explain how this works later on.

Fig. 12.12 Ceramic multilayer actuators consist of thin layers of piezoelectric ceramic and metal electrodes. In contrast to traditional piezoelectrics, even low voltages produce large forces and substantial displacements. A tradeoff exists between force and displacement. The multilayer stack utilizing the d_{33} coefficient give kilonewton forces capable of pushing heavy weights through small distances. Bimorph benders make use of the smaller transverse of d_{31} coefficients to give larger displacements in the millimeter range, but only small forces.

[Newnham]

Piezoelectric Crystals

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- How is it that crystals can be piezoelectric?
- The answer is that the bonding must be ionic to some degree (i.e. there is a net charge on the different elements) and the arrangement of the atoms must be non-centrosymmetric.
- PZT is a standard piezoelectric material. It has Pb atoms at the cell corners ($a \sim 4\text{\AA}$), O on face centers, and a Ti or Zr atom near the body center. Below a certain temperature (*Curie T*), the cell transforms from cubic (high T) to tetragonal (low T). Applying stress distorts the cell, which changes the electric displacement in different ways (see figure).
- Although we can understand the effect at the single crystal level, real devices (e.g. sonar transducers) are polycrystalline. The operation is much complicated than discussed here, and involves “poling” to maximize the response, which in turns involves motion of domain walls.

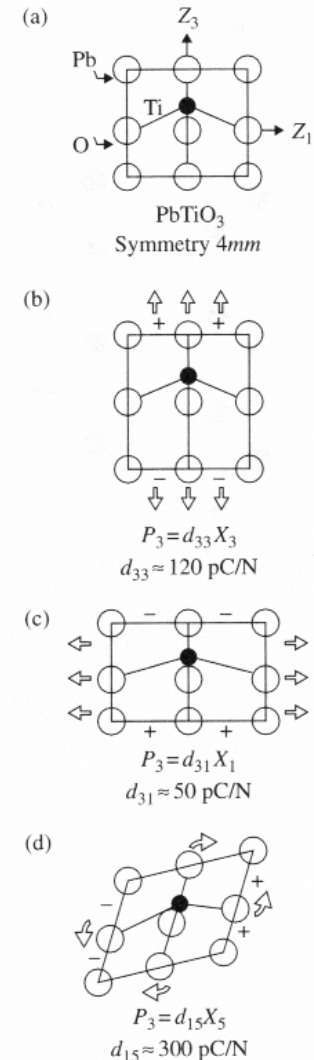


Fig. 12.11 Structure–property relations for the intrinsic piezoelectric effect in PbTiO_3 . In the unstressed state there is an electric dipole associated with the off-center shift of the titanium atom. Under stress, this dipole can be increased (d_{33}), decreased (d_{31}), or tilted (d_{15}).

[Newnham]

Mathematical Descriptions

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- Mathematical descriptions of properties are available.
- Mathematics, or a type of mathematics provides a *quantitative framework*. It is always necessary, however, to make a correspondence between mathematical variables and physical quantities.
- In group theory one might say that there is a set of mathematical operations & parameters, and a set of physical quantities and processes: if the mathematics is a good description, then the two sets are isomorphous.
- This lecture makes extensive use of *tensors*. A tensor is a quantity that can be transformed from one set of axes to another via the *tensor transformation rule* (next slide).

Tensor: definition, contd.

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- In order for a quantity to “qualify” as a *tensor* it has to obey the *axis transformation rule*, as discussed in the previous slides.
- The *transformation rule* defines relationships between transformed and untransformed tensors of various ranks.

Vector:

$$V'_i = a_{ij}V_j$$

2nd rank

$$T'_{ij} = a_{ik}a_{il}T_{kl}$$

3rd rank

$$T'_{ijk} = a_{il}a_{im}a_{kn}T_{lmn}$$

4th rank

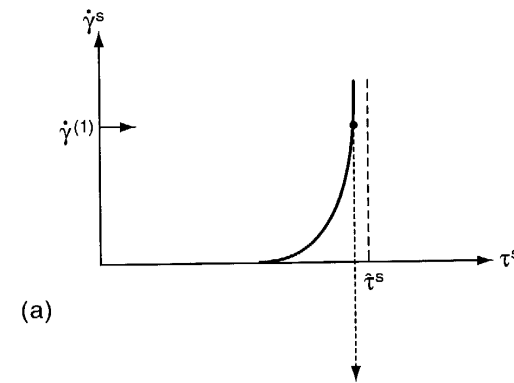
$$T'_{ijkl} = a_{im}a_{in}a_{ko}a_{lp}T_{mnop}$$

This rule is a critical piece of information, which you must know how to use.

Non-Linear properties, example

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- Another important example of non-linear anisotropic properties is plasticity, i.e. the irreversible deformation of solids.
- A typical description of the response at plastic yield (what happens when you load a material to its yield stress) is elastic-perfectly plastic. In other words, the material responds elastically until the yield stress is reached, at which point the stress remains constant (strain rate unlimited).
- A more realistic description is a power-law with a large exponent, $n \sim 50$. The stress is scaled by the *crss*, and be expressed as either shear stress-shear strain rate [graph], or tensile stress-tensile strain [equation].



$$\dot{\epsilon} = \left(\frac{\sigma}{\sigma_{yield}} \right)^n$$

[Kocks]

Linear properties

- Certain properties, such as elasticity in most cases, are linear which means that we can simplify even further to obtain

$$R = R_0 + \mathbf{P}F$$

or if $R_0 = 0$,

$$R = \mathbf{P}F.$$

e.g. elasticity: $\sigma = C \varepsilon$

stiffness



In tension, $C \equiv \text{Young's modulus, } Y \text{ or } E.$

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- *Elasticity*: example of a property that requires tensors to describe it fully.
- Even in cubic metals, a crystal is quite anisotropic. The [111] in many cubic metals is stiffer than the [100] direction.
- Even in cubic materials, 3 numbers/coefficients/moduli are required to describe elastic properties; isotropic materials only require 2.
- *Familiarity with Miller indices, suffix notation, Einstein convention, Kronecker delta, permutation tensor, and tensors is assumed.*

Elastic Anisotropy: 1

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- First we restate the linear elastic relations for the properties *Compliance*, written S , and *Stiffness*, written C (admittedly not very logical choice of notation), which connect stress, σ , and strain, ε . We write it first in vector-tensor notation with “:” signifying *inner product* (i.e. add up terms that have a common suffix or index in them):

$$\sigma = C:\varepsilon$$

$$\varepsilon = S:\sigma$$

- In *component* form (with suffices),

$$\sigma_{ij} = C_{ijkl}\varepsilon_{kl}$$

$$\varepsilon_{ij} = S_{ijkl}\sigma_{kl}$$

Elastic Anisotropy: 2

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The definitions of the stress and strain tensors mean that they are both symmetric (second rank) tensors. Therefore we can see that

$$\varepsilon_{23} = S_{2311} \sigma_{11}$$

$$\varepsilon_{32} = S_{3211} \sigma_{11} = \varepsilon_{23}$$

which means that,

$$S_{2311} = S_{3211}$$

and in general,

$$S_{ijkl} = S_{jikl}$$

We will see later on that this reduces considerably the number of different coefficients needed.

Stiffness in sample coords.

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- Consider how to express the elastic properties of a single crystal in the sample coordinates. In this case we need to rotate the (4th rank) tensor stiffness from crystal coordinates **to** sample coordinates using the orientation (matrix), a :

$$C_{ijkl}' = a_{im} a_{jn} a_{ko} a_{lp} C_{mnop}$$

- Note how the transformation matrix appears four times because we are transforming a 4th rank tensor!
- The axis transformation matrix, a , is sometimes also written as λ , also as the transpose of the **orientation matrix** g^T .

Young's modulus from compliance

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- Young's modulus as a function of direction can be obtained from the compliance tensor as:

$$E = 1/s'_{1111}$$

Using compliances and a stress boundary condition (only $\sigma_{11} \neq 0$) is most straightforward. To obtain s'_{1111} , we simply apply the same transformation rule,

$$s'_{ijkl} = a_{im} a_{jn} a_{ko} a_{lp} s_{mnop}$$

“Voigt” or “matrix” notation

- It is useful to re-express the three quantities involved in a simpler format. The stress and strain tensors are **vectorized**, i.e. converted into a 1x6 notation and the elastic tensors are reduced to 6x6 matrices.

$$\begin{pmatrix} \sigma_{11} & \sigma_{12} & \sigma_{13} \\ \sigma_{21} & \sigma_{22} & \sigma_{23} \\ \sigma_{31} & \sigma_{32} & \sigma_{33} \end{pmatrix} \longleftrightarrow \begin{pmatrix} \sigma_1 & \sigma_6 & \sigma_5 \\ \sigma_6 & \sigma_2 & \sigma_4 \\ \sigma_5 & \sigma_4 & \sigma_3 \end{pmatrix} \\
 \longleftrightarrow (\sigma_1, \sigma_2, \sigma_3, \sigma_4, \sigma_5, \sigma_6)$$

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“matrix notation”, contd.

- Similarly for strain:

$$\begin{pmatrix} \varepsilon_{11} & \varepsilon_{12} & \varepsilon_{13} \\ \varepsilon_{21} & \varepsilon_{22} & \varepsilon_{23} \\ \varepsilon_{31} & \varepsilon_{32} & \varepsilon_{33} \end{pmatrix} \longleftrightarrow \begin{pmatrix} \varepsilon_1 & \frac{1}{2}\varepsilon_6 & \frac{1}{2}\varepsilon_5 \\ \frac{1}{2}\varepsilon_6 & \varepsilon_2 & \frac{1}{2}\varepsilon_4 \\ \frac{1}{2}\varepsilon_5 & \frac{1}{2}\varepsilon_4 & \varepsilon_3 \end{pmatrix}$$

$$\longleftrightarrow (\varepsilon_1, \varepsilon_2, \varepsilon_3, \varepsilon_4, \varepsilon_5, \varepsilon_6)$$

The particular definition of shear strain used in the reduced notation happens to correspond to that used in mechanical engineering such that ε_4 is the change in angle between direction 2 and direction 3 due to deformation.

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Work conjugacy, matrix inversion

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- The more important consideration is that the reason for the factors of two is so that work conjugacy is maintained.

$$dW = \sigma : d\varepsilon = \sigma_{ij} : d\varepsilon_{ij} = \sigma_k \bullet d\varepsilon_k$$

Also we can combine the expressions

$$\sigma = C\varepsilon \text{ and } \varepsilon = S\sigma \text{ to give:}$$

$$\sigma = CS\sigma, \text{ which shows:}$$

$$I = CS, \text{ or, } C = S^{-1}$$

Stiffness Matrix

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$$\mathbf{C} = \begin{bmatrix} C_{11} & C_{12} & C_{13} & C_{14} & C_{15} & C_{16} \\ C_{21} & C_{22} & C_{23} & C_{24} & C_{25} & C_{26} \\ C_{31} & C_{32} & C_{33} & C_{34} & C_{35} & C_{36} \\ C_{41} & C_{42} & C_{43} & C_{44} & C_{45} & C_{46} \\ C_{51} & C_{52} & C_{53} & C_{54} & C_{55} & C_{56} \\ C_{61} & C_{62} & C_{63} & C_{64} & C_{65} & C_{66} \end{bmatrix}$$

Tensor conversions: compliance

- For compliance some factors of two are required and so the rule becomes:

$$p S_{ijkl} = S_{mn}$$

$$p = 1$$

$$m.AND.n \in [1,2,3]$$

$$p = 2$$

$$m.XOR.n \in [1,2,3]$$

$$p = 4$$

$$m.AND.n \in [4,5,6]$$

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Relationships between coefficients: C in terms of S

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Some additional useful relations between coefficients for cubic materials are as follows. Symmetrical relationships exist for compliances in terms of stiffnesses (next slide).

$$C_{11} = (S_{11} + S_{12}) / \{(S_{11} - S_{12})(S_{11} + 2S_{12})\}$$

$$C_{12} = -S_{12} / \{(S_{11} - S_{12})(S_{11} + 2S_{12})\}$$

$$C_{44} = 1/S_{44}.$$

S in terms of C

The relationships for S in terms of C are symmetrical to those for stiffnesses in terms of compliances (a simple exercise in algebra).

$$S_{11} = (C_{11} + C_{12}) / \{(C_{11} - C_{12})(C_{11} + 2C_{12})\}$$

$$S_{12} = -C_{12} / \{(C_{11} - C_{12})(C_{11} + 2C_{12})\}$$

$$S_{44} = 1/C_{44}.$$

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Neumann's Principle

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- A fundamental natural law: *Neumann's Principle: the symmetry elements of any physical property of a crystal must include the symmetry elements of the point group of the crystal.* The property may have additional symmetry elements to those of the crystal (point group) symmetry. There are 32 crystal classes for the point group symmetry.
- F.E. Neumann 1885.

Neumann, extended

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- If a crystal has a defect structure such as a dislocation network that is arranged in a non-uniform way then the symmetry of certain properties may be reduced from the crystal symmetry. In principle, a finite elastic strain in one direction decreases the symmetry of a cubic crystal to tetragonal or less. Therefore the modified version of *Neumann's Principle: the symmetry elements of any physical property of a crystal must include the symmetry elements that are common to the point group of the crystal and the defect structure contained within the crystal.*

Effect of crystal symmetry

- Consider an active rotation of the crystal, where O is the symmetry operator. Since the crystal is indistinguishable (looks the same) after applying the symmetry operator, the result before, $R^{(1)}$, and the result after, $R^{(2)}$, must be identical:

$$\left. \begin{aligned} R^{(1)} &= \mathbf{P}F \\ R^{(2)} &= \mathbf{O}\mathbf{P}\mathbf{O}^T F \\ R^{(1)} &\overset{=}{\longleftrightarrow} R^{(2)} \end{aligned} \right\}$$

The two results are indistinguishable and therefore equal. It is essential, however, to express the property and the operator in the same (crystal) reference frame.

Symmetry, properties, contd.

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- Expressed mathematically, we can rotate, e.g. a second rank property tensor thus:

$$\mathbf{P}' = \mathbf{O}\mathbf{P}\mathbf{O}^T = \mathbf{P}, \text{ or, in coefficient notation,}$$

$$P'_{ij} = O_{ik}O_{il}P_{kl}$$

where O is a symmetry operator.

- Since the rotated (property) tensor, \mathbf{P}' , must be the same as the original tensor, \mathbf{P} , then we can equate coefficients:

$$P'_{ij} = P_{ij}$$

- If we find, for example, that $P'_{21} = -P_{21}$, then the only value of P_{21} that satisfies this equality is $P_{21} = 0$.
- Remember that you must express the property with respect to a particular set of axes in order to use the coefficient form. In everything related to single crystals, *always* use the crystal axes as the reference frame!
- Homework question: based on cubic crystal symmetry, work out why a second rank tensor property can only have *one* independent coefficient.

Effect of symmetry on stiffness matrix

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- Why do we need to look at the effect of symmetry? For a cubic material, only 3 *independent* coefficients are needed as opposed to the 81 coefficients in a 4th rank tensor. The reason for this is the symmetry of the material.
- What does symmetry mean? Fundamentally, if you pick up a crystal, rotate [mirror] it and put it back down, then a symmetry operation [rotation, mirror] is such that *you cannot tell that anything happened*.
- From a mathematical point of view, this means that the property (its coefficients) *does not change*. For example, if the symmetry operator changes the sign of a coefficient, then it must be equal to zero.

2nd Rank Tensor Properties & Symmetry

TABLE 3

The effect of crystal symmetry on properties represented by symmetrical second-rank tensors

Optical classification	System	Characteristic symmetry (see p. 280)†	Nature of representation quadric and its orientation	Number of independent coefficients	Tensor referred to axes in the conventional orientation‡
Isotropic (anaxial)	Cubic	4 3-fold axes	Sphere	1	$\begin{bmatrix} S & 0 & 0 \\ 0 & S & 0 \\ 0 & 0 & S \end{bmatrix}$
Uniaxial	Tetragonal	1 4-fold axis	Quadric of revolution about the principal symmetry axis (x_3)(z)	2	$\begin{bmatrix} S_1 & 0 & 0 \\ 0 & S_1 & 0 \\ 0 & 0 & S_3 \end{bmatrix}$
	Hexagonal	1 6-fold axis			
	Trigonal	1 3-fold axis			
Biaxial	Orthorhombic	3 mutually perpendicular 2-fold axes; no axes of higher order	General quadric with axes (x_1, x_2, x_3) to the diad axes (x, y, z)	3	$\begin{bmatrix} S_1 & 0 & 0 \\ 0 & S_2 & 0 \\ 0 & 0 & S_3 \end{bmatrix}$
	Monoclinic	1 2-fold axis	General quadric with one axis (x_2) to the diad axis (y)	4	$\begin{bmatrix} S_{11} & 0 & S_{31} \\ 0 & S_2 & 0 \\ S_{31} & 0 & S_{33} \end{bmatrix}$
	Triclinic	A centre of symmetry or no symmetry	General quadric. No fixed relation to crystallographic axes	6	$\begin{bmatrix} S_{11} & S_{12} & S_{31} \\ S_{12} & S_{22} & S_{23} \\ S_{31} & S_{23} & S_{33} \end{bmatrix}$

† Axes of symmetry may be rotation axes or inversion axes.

‡ The setting of the reference axes x_1, x_2, x_3 in column 6 in relation to the crystallographic axes x, y, z and to the symmetry elements is that shown in column 4. For further notes on axial conventions, see Appendix B.

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- The table from Nye shows the number of independent, non-zero coefficients allowed in a 2nd rank tensor according to the crystal symmetry class.

Effect of symmetry on stiffness matrix

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- Following Reid, p.66 *et seq.*:

Apply a 90° rotation about the crystal-z axis (axis 3),

$$C'_{ijkl} = O_{im}O_{jn}O_{ko}O_{lp}C_{mnop}:$$

$$C' = C$$

$$O_4^z = \begin{pmatrix} 0 & -1 & 0 \\ 1 & 0 & 0 \\ 0 & 0 & 1 \end{pmatrix}$$

$$C' = \begin{bmatrix} C_{22} & C_{21} & C_{23} & C_{25} & -C_{24} & -C_{26} \\ C_{21} & C_{11} & C_{13} & C_{15} & -C_{14} & -C_{16} \\ C_{23} & C_{13} & C_{33} & C_{35} & -C_{34} & -C_{36} \\ C_{25} & C_{15} & C_{35} & C_{55} & -C_{54} & -C_{56} \\ -C_{24} & -C_{14} & -C_{34} & -C_{54} & C_{44} & C_{46} \\ -C_{26} & -C_{16} & -C_{36} & -C_{56} & C_{46} & C_{66} \end{bmatrix}$$

Effect of symmetry, 2

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- Using $P' = P$, we can equate coefficients and find that:

$$C_{11} = C_{22}, \quad C_{13} = C_{23}, \quad C_{44} = C_{35}, \quad C_{16} = -C_{26},$$

$$C_{14} = C_{15} = C_{24} = C_{25} = C_{34} = C_{35} = C_{36} = C_{45} = C_{46} =$$

$$C_{56} = 0.$$

$$C' = \begin{bmatrix} C_{11} & C_{12} & C_{13} & 0 & 0 & C_{16} \\ C_{12} & C_{11} & C_{13} & 0 & 0 & -C_{16} \\ C_{13} & C_{13} & C_{33} & 0 & 0 & 0 \\ 0 & 0 & 0 & C_{44} & 0 & 0 \\ 0 & 0 & 0 & 0 & C_{44} & C_{46} \\ C_{16} & -C_{16} & 0 & 0 & C_{46} & C_{66} \end{bmatrix}$$

Effect of symmetry, 3

- Thus by repeated applications of the symmetry operators, one can demonstrate (for cubic crystal symmetry) that one can reduce the 81 coefficients down to only 3 independent quantities. These become two in the case of isotropy.

$$\begin{bmatrix} C_{11} & C_{12} & C_{12} & 0 & 0 & 0 \\ C_{12} & C_{11} & C_{12} & 0 & 0 & 0 \\ C_{12} & C_{12} & C_{11} & 0 & 0 & 0 \\ 0 & 0 & 0 & C_{44} & 0 & 0 \\ 0 & 0 & 0 & 0 & C_{44} & 0 \\ 0 & 0 & 0 & 0 & 0 & C_{44} \end{bmatrix}$$

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Cubic crystals: anisotropy factor

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- If one applies the symmetry elements of the cubic system, it turns out that only three independent coefficients remain: C_{11} , C_{12} and C_{44} , (similar set for compliance). From these three, a useful combination of the first two is

$$C' = (C_{11} - C_{12})/2$$

- See Nye, *Physical Properties of Crystals*

Zener's anisotropy factor

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- $C' = (C_{11} - C_{12})/2$ turns out to be the stiffness associated with a shear in a $\langle 110 \rangle$ direction on a plane. In certain martensitic transformations, this modulus can approach zero which corresponds to a structural instability. Zener proposed a measure of elastic anisotropy based on the ratio C_{44}/C' . This turns out to be a useful criterion for identifying materials that are elastically anisotropic.

Rotated compliance (matrix)

Objective

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- Given an orientation a_{ij} , we transform the compliance tensor, using cubic point group symmetry, and find that:

$$\begin{aligned}
 S'_{11} = & S_{11} \left(a_{11}^4 + a_{12}^4 + a_{13}^4 \right) \\
 & + 2S_{12} \left(a_{12}^2 a_{13}^2 + a_{11}^2 a_{12}^2 + a_{11}^2 a_{13}^2 \right) \\
 & + S_{44} \left(a_{12}^2 a_{13}^2 + a_{11}^2 a_{12}^2 + a_{11}^2 a_{13}^2 \right)
 \end{aligned}$$

Rotated compliance (matrix)

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- This can be further simplified with the aid of the standard relations between the direction cosines, $a_{ik}a_{jk} = 1$ for $i=j$; $a_{ik}a_{jk} = 0$ for $i \neq j$, ($a_{ik}a_{jk} = \delta_{ij}$) to read as follows.

$$s'_{11} = s_{11} -$$

$$2 \left(s_{11} - s_{12} - \frac{s_{44}}{2} \right) \left\{ \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right\}$$

- By definition, the Young's modulus in any direction is given by the reciprocal of the compliance, $E = 1/S'_{11}$.

Anisotropy in cubic materials

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- Thus the second term on the RHS is zero for $\langle 100 \rangle$ directions and, for $C_{44}/C' > 1$, a maximum in $\langle 111 \rangle$ directions (conversely a minimum for $C_{44}/C' < 1$). The following table shows that most cubic metals have positive values of Zener's coefficient so that $\langle 100 \rangle$ is soft and $\langle 111 \rangle$ is hard, with the exceptions of V and NaCl.

Material	C_{44}/C'	E_{111}/E_{100}
Cu	3.21	2.87
Ni	2.45	2.18
Al	1.22	1.19
Fe	2.41	2.15
Ta	1.57	1.50
W (2000K)	1.23	1.35
W (R.T.)	1.01	1.01
V	0.78	0.72
Nb	0.55	0.57
β -CuZn	18.68	8.21
spinel	2.43	2.13
MgO	1.49	1.37
NaCl	0.69	0.74

Stiffness coefficients, cubics

Material class	Material	C_{11} (10^{10} N/m ²)	C_{12} (10^{10} N/m ²)	C_{44} (10^{10} N/m ²)	Anisotropy ratio ($C_{11} - C_{12}$)/ $2C_{44}$
Metals	Ag	12.4	9.3	4.6	0.34
	Al	10.8	6.1	2.9	0.81
	Au	18.6	15.7	4.2	0.35
	Cu	16.8	12.1	7.5	0.31
	α -Fe	23.7	14.1	11.6	0.41
	Mo	46.0	17.6	11.0	1.29
	Na	0.73	0.63	0.42	0.12
	Ni	24.7	14.7	12.5	0.40
	Pb	5.0	4.2	1.5	0.27
	W	50.1	19.8	15.1	1.00
Covalent solids	Si	16.6	6.4	8.0	0.64
	Diamond	107.6	12.5	57.6	0.83
	TiC	51.2	11.0	17.7	1.14
Ionic solids	LiF	11.2	4.6	6.3	0.52
	MgO	29.1	9.0	15.5	0.65
	NaCl	4.9	1.3	1.3	1.38

Table 2.2
Stiffness coefficients for selected cubic materials

[Courtney]

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Anisotropy in terms of moduli

- Another way to write the above equation is to insert the values for the Young's modulus in the soft and hard directions, assuming that the $\langle 100 \rangle$ are the most compliant direction(s). (Courtney uses α , β , and γ in place of my α_1 , α_2 , and α_3 .) The advantage of this formula is that moduli in specific directions can be used directly.

$$\frac{1}{E_{uvw}} = \frac{1}{E_{100}} - 3 \left\{ \frac{1}{E_{100}} - \frac{1}{E_{111}} \right\} \left(\alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right)$$

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Example Problem

2.11 a Sketch a (001) plane in a face-centered cubic material and an arbitrary vector within it making an angle θ with the [100] direction. Plot the Young's modulus for copper as a function of θ for directions between [110] and [100].
 b Sketch a (110) plane in Cu and a vector in the plane making an angle α with the [110] direction. Plot E vs. α for directions between [110] and [001].

Solution: (a) The plane is illustrated to the right. Use Eq. (3.22).

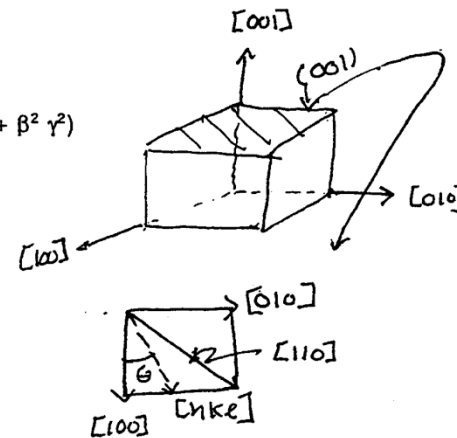
$$1/E_{[hkl]} = 1/E_{\langle 100 \rangle} - 3\{1/E_{\langle 100 \rangle} - 1/E_{\langle 111 \rangle}\}(\alpha^2\beta^2 + \alpha^2\gamma^2 + \beta^2\gamma^2)$$

where α is the cosine of the angle between the direction [100] and [hkl], β is the cosine of the angle between [010] and [hkl], and γ is the like cosine between [001] and [hkl]. From the sketches provided we see that $\alpha = \cos\theta$, $\beta = \cos(90^\circ - \theta) = \sin\theta$ and $\gamma = 0$. Employing moduli in units of 10^{10} N/m², with $E_{\langle 100 \rangle} = 6.7$, $E_{\langle 111 \rangle} = 11.2$ the above equation becomes

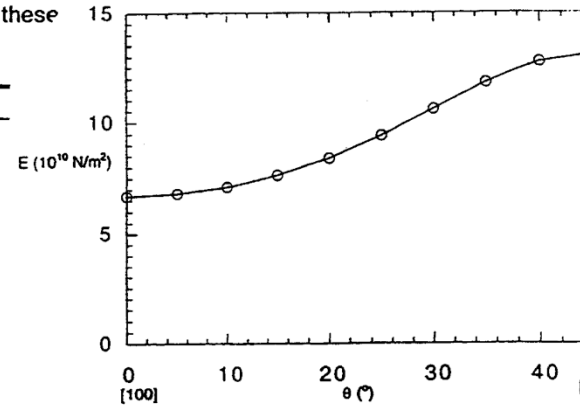
$$1/E_{[hkl]} = 0.149 - 0.2915\cos^2\theta\sin^2\theta$$

The table below presents results obtained with the above formula; the figure to the right graphs these results.

θ (°)	$\cos^2\theta\sin^2\theta$	E (10^{10} N/m ²)
0	0	6.7
5	0.0075	6.81
10	0.0292	7.12
15	0.0625	7.65
20	0.1033	8.41
25	0.1467	9.41
30	0.1875	10.6
35	0.2207	11.82
40	0.2425	12.77
45 (= [110])	0.25	13.14



Should be $E_{\langle 111 \rangle} = 18.89$



[Courtney]

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Alternate Vectorization

$$\mathbf{b}^{(1)} = \frac{1}{\sqrt{6}} \begin{pmatrix} -1 & 0 & 0 \\ 0 & -1 & 0 \\ 0 & 0 & 2 \end{pmatrix}; \quad \mathbf{b}^{(2)} = \frac{1}{\sqrt{2}} \begin{pmatrix} -1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 0 \end{pmatrix}; \quad \mathbf{b}^{(3)} = \frac{1}{\sqrt{2}} \begin{pmatrix} 0 & 0 & 0 \\ 0 & 0 & 1 \\ 0 & 1 & 0 \end{pmatrix}$$

$$\mathbf{b}^{(4)} = \frac{1}{\sqrt{2}} \begin{pmatrix} 0 & 0 & 1 \\ 0 & 0 & 0 \\ 1 & 0 & 0 \end{pmatrix}; \quad \mathbf{b}^{(5)} = \frac{1}{\sqrt{2}} \begin{pmatrix} 0 & 1 & 0 \\ 1 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix}; \quad \mathbf{b}^{(6)} = \frac{1}{\sqrt{3}} \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}$$

An alternate vectorization, discussed by Tomé on p287 of the Kocks et al. textbook, is to use the above set of eigentensors. For both stress and strain, one can matrix multiply each eigentensor into the stress/strain tensor in turn and obtain the coefficient of the corresponding stress/strain vector. Work conjugacy is still satisfied. The first two eigentensors represent shears in the $\{110\}$ planes; the next three are simple shears on $\{110\}\langle 110\rangle$ systems, and the last (6th) is the hydrostatic component. The same vectorization can be used for plastic anisotropy, except in this case, the sixth, hydrostatic component is (generally) ignored.

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Summary

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- We have covered the following topics:
 - Linear properties
 - Non-linear properties
 - Examples of properties
 - Tensors, vectors, scalars, tensor transformation law.
 - Elasticity, as example as of higher order property, also as example as how to apply (crystal) symmetry.

Supplemental Slides

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- The following slides contain some useful material for those who are not familiar with all the detailed mathematical methods of matrices, transformation of axes, tensors etc.

Einstein Convention

- The Einstein Convention, or summation rule for suffixes looks like this:

$$A_i = B_{ij} C_j$$

where “i” and “j” both are integer indexes whose range is {1,2,3}. So, to find each “ith” component of A on the LHS, we sum up over the **repeated index, “j”**, on the RHS:

$$A_1 = B_{11}C_1 + B_{12}C_2 + B_{13}C_3$$

$$A_2 = B_{21}C_1 + B_{22}C_2 + B_{23}C_3$$

$$A_3 = B_{31}C_1 + B_{32}C_2 + B_{33}C_3$$

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Matrix Multiplication

- Take each row of the LH matrix in turn and multiply it into each column of the RH matrix.
- In suffix notation, $a_{ij} = b_{ik}c_{kj}$

$$\begin{bmatrix} a\alpha + b\delta + c\gamma & a\beta + b\varepsilon + c\mu & a\gamma + b\phi + c\nu \\ d\alpha + e\delta + f\gamma & d\beta + e\varepsilon + f\mu & d\gamma + e\phi + f\nu \\ l\alpha + m\delta + n\gamma & l\beta + m\varepsilon + n\mu & l\gamma + m\phi + n\nu \end{bmatrix}$$

$$= \begin{bmatrix} a & b & c \\ d & e & f \\ l & m & n \end{bmatrix} \times \begin{bmatrix} \alpha & \beta & \gamma \\ \delta & \varepsilon & \phi \\ \lambda & \mu & \nu \end{bmatrix}$$

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Properties of Rotation Matrix

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- The rotation matrix is an *orthogonal matrix*, meaning that any row is orthogonal to any other row (the dot products are zero). In physical terms, each row represents a unit vector that is the position of the corresponding (new) old axis in terms of the (old) new axes.
- The same applies to columns: in suffix notation -

$$a_{ij}a_{kj} = \delta_{ik}, \quad a_{ji}a_{jk} = \delta_{ik}$$

The diagram shows a 3x3 matrix with elements a, b, c in the first row, d, e, f in the second row, and l, m, n in the third row. A red bracket groups the first two rows, with the equation $ad+be+cf = 0$ written to its right. A blue bracket groups the last two columns, with the equation $bc+ef+mn = 0$ written below it.

Direction Cosines, contd.

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- That the set of direction cosines are not independent is evident from the following construction:

$$\hat{e}'_i \cdot \hat{e}'_j = a_{ik} a_{jl} \hat{e}_k \cdot \hat{e}_l = a_{ik} a_{jl} \delta_{kl} = a_{ik} a_{jk} = \delta_{ij}$$

Thus, there are *six* relationships (*i* takes values from 1 to 3, and *j* takes values from 1 to 3) between the *nine* direction cosines, and therefore, as stated above, only *three* are independent, exactly as expected for a rotation.

- Another way to look at a rotation: combine an axis (described by a unit vector with two parameters) and a rotation angle (one more parameter, for a total of 3).

Orthogonal Matrices

- Note that the direction cosines can be arranged into a 3×3 matrix, Λ , and therefore the relation above is equivalent to the expression

$$\Lambda \Lambda^T = \mathbf{I}$$

where Λ^T denotes the transpose of Λ . This relationship identifies Λ as an orthogonal matrix, which has the properties

$$\Lambda^{-1} = \Lambda^T \quad \det \Lambda = \pm 1$$

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- When both coordinate systems are right-handed, $\det(\Lambda) = +1$ and Λ is a *proper orthogonal matrix*. The orthogonality of Λ also insures that, in addition to the relation above, the following holds:

$$\hat{e}_j = a_{ij} \hat{e}'_i$$

Combining these relations leads to the following inter-relationships between components of vectors in the two coordinate systems:

$$v_i = a_{ji} v'_j, \quad v'_j = a_{ji} v_i$$

Transformation Law

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- These relations are called the *laws of transformation* for the components of vectors. They are a consequence of, and equivalent to, the parallelogram law for addition of vectors. That such is the case is evident when one considers the scalar product expressed in two coordinate systems:

$$\vec{u} \cdot \vec{v} = u_i v_i = a_{ji} u'_j a_{ki} v'_k =$$

$$\delta_{jk} u'_j v'_k = u'_j v'_j = u'_i v'_i$$

Invariants

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Thus, the transformation law as expressed preserves the lengths and the angles between vectors. Any function of the components of vectors which remains unchanged upon changing the coordinate system is called an *invariant* of the vectors from which the components are obtained. The derivations illustrate the fact that the scalar product $\vec{u} \cdot \vec{v}$ is an *invariant* of \vec{u} and \vec{v} . Other examples of *invariants* include the vector product of two vectors and the triple scalar product of three vectors. The reader should note that the transformation law for vectors also applies to the components of points when they are referred to a common origin.

Orthogonality

- A rotation matrix, Λ , is an *orthogonal matrix*, however, because each row is mutually orthogonal to the other two.

$$a_{ki}a_{kj} = \delta_{ij}, \quad a_{ik}a_{jk} = \delta_{ij}$$

- Equally, each column is orthogonal to the other two, which is apparent from the fact that each row/column contains the direction cosines of the new/old axes in terms of the old/new axes and we are working with [mutually perpendicular] Cartesian axes.

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Anisotropy

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- *Anisotropy* as a word simply means that something varies with direction.
- Anisotropy is from the Greek: *aniso* = different, varying; *tropos* = direction.
- Almost all crystalline materials are anisotropic; many materials are engineered to take advantage of their anisotropy (beer cans, turbine blades, microchips...)
- Older texts use trigonometric functions to describe anisotropy but tensors offer a general description with which it is much easier to perform calculations.
- For materials, what we know is that some properties are anisotropic. This means that several numbers, or *coefficients*, are needed to describe the property - one number is not sufficient.
- Elasticity is an important example of a property that, when examined in single crystals, is often highly anisotropic. In fact, the lower the crystal symmetry, the greater the anisotropy is likely to be.
- *Nomenclature*: in general, we need to use *tensors* to describe fields and properties. The simplest case of a tensor is a *scalar* which is all we need for isotropic properties. The next “level” of tensor is a *vector*, e.g. electric current.

Scalars, Vectors, Tensors

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- *Scalar*:= quantity that requires only one number, e.g. density, mass, specific heat. Equivalent to a zero-rank tensor.
- *Vector*:= quantity that has direction as well as magnitude, e.g. velocity, current, magnetization; requires 3 numbers or *coefficients* (in 3D). Equivalent to a first-rank tensor.
- *Tensor*:= quantity that requires higher order descriptions but is the same, no matter what coordinate system is used to describe it, e.g. stress, strain, elastic modulus; requires 9 (or more, depending on rank) numbers or *coefficients*.

Vector field, response

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- If we have a vector response, R , that we can write in component form, a vector field, F , that we can also write in component form, and a property, P , that we can write in matrix form (with nine coefficients) then the linearity of the property means that we can write the following ($R_0 = 0$):

$$R_i = P_{ij}F_j$$

- A *scalar* (e.g. pressure) can be called a *zero-rank tensor*.
- A *vector* (e.g. electric current) is also known as a *first-rank tensor*.

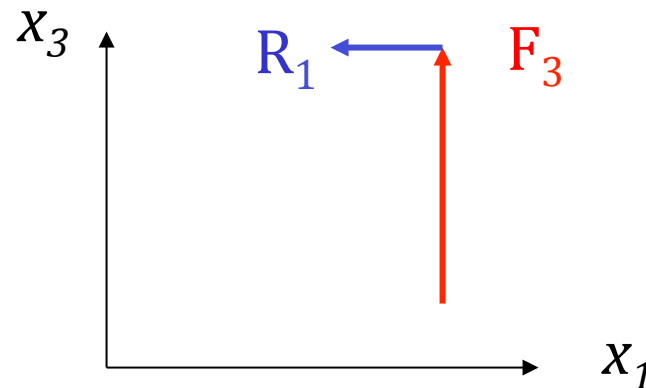
Linear anisotropic property

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- This means that each component of the response is *linearly* related to each component of the field and that the proportionality constant is the appropriate coefficient in the matrix. Example:

$$R_1 = P_{13}F_3,$$

which says that the first component of the response is linearly related to the third field component through the property coefficient P_{13} .



Example: electrical conductivity

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- An example of such a linear anisotropic (second order tensor, discussed in later slides) property is the electrical conductivity of a material:
 - Field: Electric Field, E
 - Response: Current Density, J
 - Property: Conductivity, σ
 - $J_i = \sigma_{ij} E_j$

Anisotropic electrical conductivity

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- We can illustrate anisotropy with Nye's example of electrical conductivity, σ :

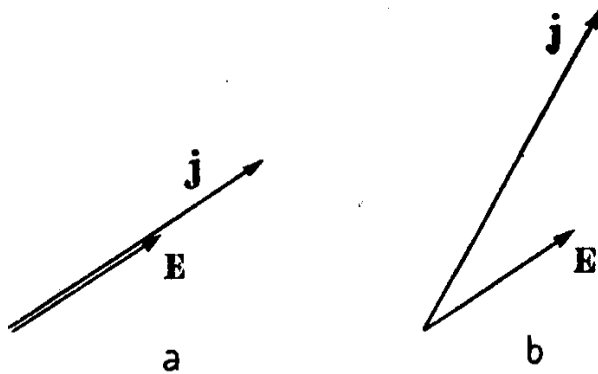


FIG. 1.1. The relation between the electric current density \mathbf{j} and the electric field \mathbf{E} in (a) an isotropic conductor and (b) an anisotropic conductor.

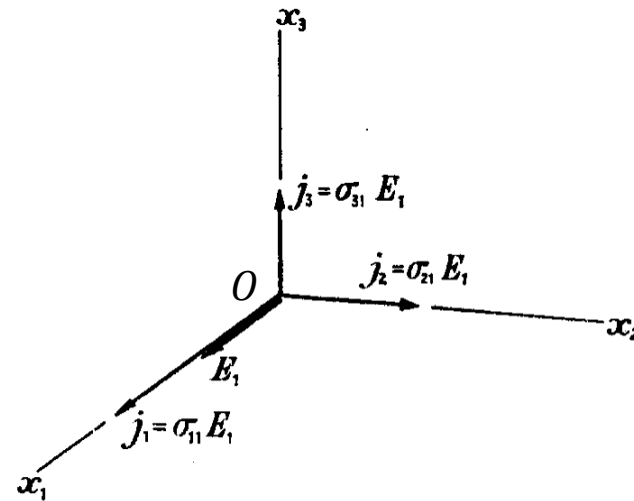


FIG. 1.2. The components of current density when a field is applied along Ox_1 .

Stimulus/ Field: $E_1 \neq 0, E_2 = E_3 = 0$

Response: $j_1 = \sigma_{11} E_1, j_2 = \sigma_{21} E_1, j_3 = \sigma_{31} E_1,$

Changing the Coordinate System

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- Many different choices are possible for the orthonormal base vectors and origin of the Cartesian coordinate system. A vector is an example of an entity which is independent of the choice of coordinate system. Its direction and magnitude must not change (and are, in fact, invariants), although its components will change with this choice.
- Why would we want to do something like this? For example, although the *properties* are conveniently expressed in a crystal reference frame, experiments often place the crystals in a non-symmetric position with respect to an experimental frame. Therefore we need some way of converting the coefficients of the property into the experimental frame.
- Changing the coordinate system is also known as *axis transformation*.

Tensor: definition, contd.

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- In order for a quantity to “qualify” as a *tensor* it has to obey the *axis transformation rule*, as discussed in the previous slides.
- The *transformation rule* defines relationships between transformed and untransformed tensors of various ranks.

Vector:

2nd rank3rd rank4th rank

$$V'_i = a_{ij}V_j$$

$$T'_{ij} = a_{ik}a_{il}T_{kl}$$

$$T'_{ijk} = a_{il}a_{im}a_{kn}T_{lmn}$$

$$T'_{ijkl} = a_{im}a_{in}a_{ko}a_{lp}T_{mnop}$$

- **This rule is a critical piece of information, which you must know how to use.**

Motivation for Axis Transformation

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- One motivation for axis transformations is the need to solve problems where the specimen shape (and the stimulus direction) does not align with the crystal axes. Consider what happens when you apply a force parallel to the sides of this specimen ...

The direction parallel to the long edge does not line up with any simple, low index crystal direction. Therefore we have to find a way to *transform* the properties that we know for the material into the frame of the problem (or vice versa).

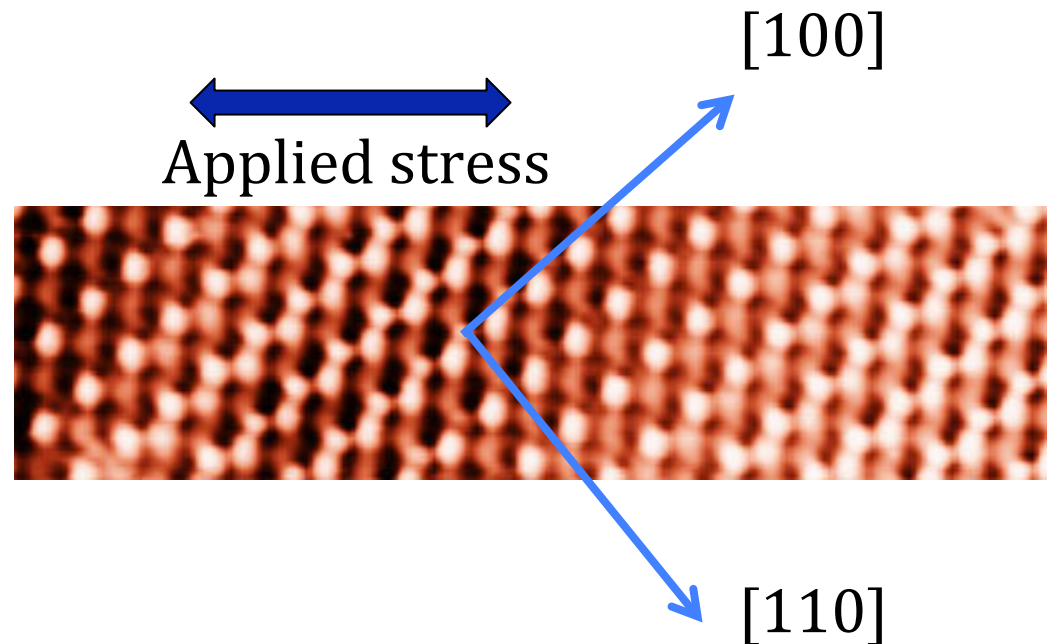


Image of Pt surface from www.cup.uni-muenchen.de/pc/wintterlin/IMGs/pt10p3.jpg

New Axes

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- Consider a *new* orthonormal system consisting of right-handed base vectors: \hat{e}'_1, \hat{e}'_2 and \hat{e}'_3
These all have the same origin, o ,
associated with \hat{e}'_1, \hat{e}'_2 and \hat{e}'_3

- The vector v is clearly expressed equally well in either coordinate system:

$$\vec{v} = v_i \hat{e}_i = v'_i \hat{e}'_i$$

Note - same physical vector but different values of the components.

- We need to find a relationship between the two sets of components for the vector.

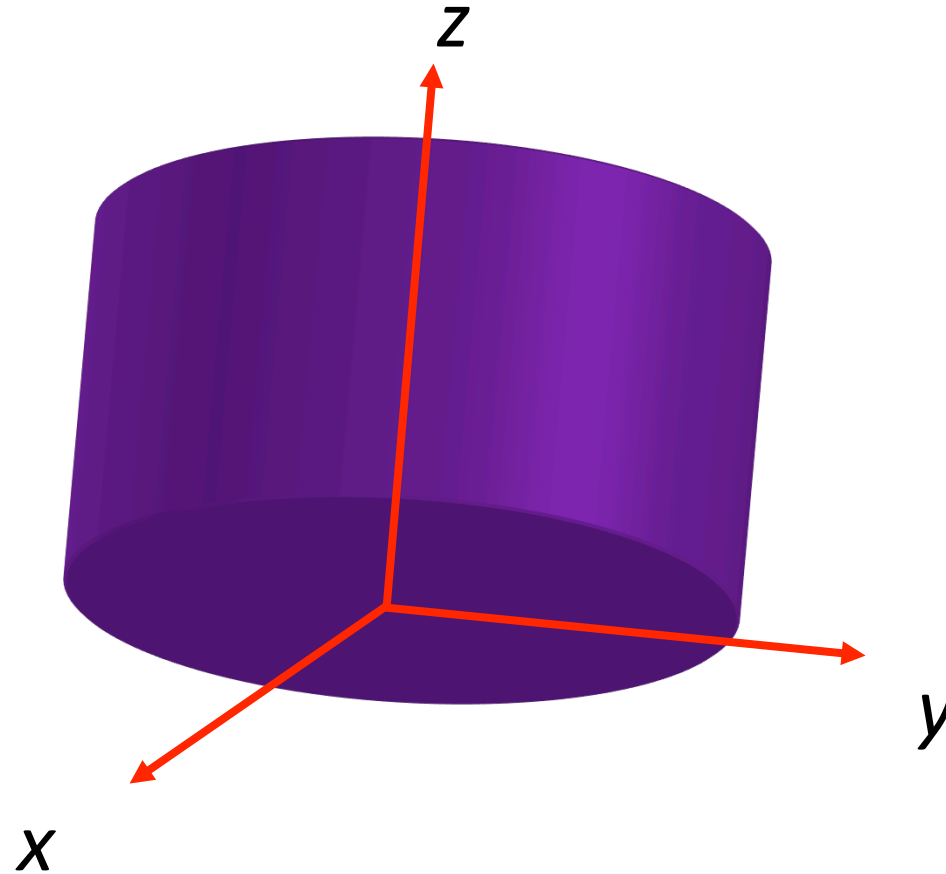
Anisotropy in Composites

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- The same methods developed here for describing the anisotropy of single crystals can be applied to composites.
- Anisotropy is important in composites, *not* because of the intrinsic properties of the components but because of the *arrangement* of the components.
- As an example, consider (a) a uniaxial composite (e.g. tennis racket handle) and (b) a flat panel cross-ply composite (e.g. wing surface).

Fiber Symmetry

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Fiber Symmetry

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- We will use the same *matrix notation* for stress, strain, stiffness and compliance as for single crystals.
- The compliance matrix, \mathbf{s} , has 5 independent coefficients.

$$\begin{bmatrix} s_{11} & s_{12} & s_{13} & 0 & 0 & 0 \\ s_{12} & s_{11} & s_{13} & 0 & 0 & 0 \\ s_{13} & s_{13} & s_{33} & 0 & 0 & 0 \\ 0 & 0 & 0 & s_{44} & 0 & 0 \\ 0 & 0 & 0 & 0 & s_{44} & 0 \\ 0 & 0 & 0 & 0 & 0 & 2(s_{11} - s_{12}) \end{bmatrix}$$

Relationships

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- For a uniaxial stress along the z (3) direction,

$$E_3 = \frac{\sigma_3}{\varepsilon_3} = \frac{1}{s_{33}} \left(= \frac{\sigma_{zz}}{\varepsilon_{zz}} \right)$$

- This stress causes strain in the transverse plane: $e_{11} = e_{22} = s_{12}\sigma_{33}$. Therefore we can calculate Poisson's ratio as:

$$\nu_{13} = \frac{e_1}{e_3} = \frac{s_{13}}{s_{33}} \left(= \frac{e_{xx}}{e_{zz}} \right)$$

- Similarly, stresses applied perpendicular to z give rise to different moduli and Poisson's ratios.

$$E_1 = \frac{\sigma_1}{\varepsilon_1} = \frac{1}{s_{11}}, \quad \nu_{21} = \frac{-s_{12}}{s_{11}}, \quad \nu_{31} = \frac{-s_{13}}{s_{11}}$$

Relationships, contd.

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Elasticity
Symmetry

- Similarly the torsional modulus is related to shears involving the z axis, i.e. yz or xz shears:

$$s_{44} = s_{55} = 1/G$$

- Shear in the x-y plane (1-2 plane) is related to the other compliance coefficients:

$$s_{66} = 2(s_{11} - s_{12}) = 1/G_{xy}$$

Plates: Orthotropic Symmetry

Objective
Linear
Ferro-
magnets
Non-linear
properties
Electric.
Conduct.
Tensors
Elasticity
Symmetry

- Again, we use the same *matrix notation* for stress, strain, stiffness and compliance as for single crystals.
- The compliance matrix, \mathbf{s} , has *9 independent coefficients*.
- This corresponds to *orthorhombic sample symmetry*: see the following slide with Table from Nye's book.

$$\begin{bmatrix} s_{11} & s_{12} & s_{13} & 0 & 0 & 0 \\ s_{12} & s_{22} & s_{23} & 0 & 0 & 0 \\ s_{13} & s_{23} & s_{33} & 0 & 0 & 0 \\ 0 & 0 & 0 & s_{44} & 0 & 0 \\ 0 & 0 & 0 & 0 & s_{55} & 0 \\ 0 & 0 & 0 & 0 & 0 & s_{66} \end{bmatrix}$$

Plates: 0° and 90° plies

Objective
Linear
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Non-linear
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Conduct.
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Symmetry

- If the composite is a laminate composite with fibers laid in at 0° and 90° in equal thicknesses then the symmetry is higher because the x and y directions are equivalent.
- The compliance matrix, s , has *6 independent coefficients*.
- This corresponds to (tetragonal) *4mm sample symmetry*: see the following slide with Table from Nye's book.

$$\begin{bmatrix} s_{11} & s_{12} & s_{13} & 0 & 0 & 0 \\ s_{12} & s_{11} & s_{13} & 0 & 0 & 0 \\ s_{13} & s_{13} & s_{33} & 0 & 0 & 0 \\ 0 & 0 & 0 & s_{44} & 0 & 0 \\ 0 & 0 & 0 & 0 & s_{44} & 0 \\ 0 & 0 & 0 & 0 & 0 & s_{66} \end{bmatrix}$$

TABLE 9
Form of the (s_{ij}) and (c_{ij}) matrices

KEY TO NOTATION

- zero component
- non-zero component
- equal components
- components numerically equal, but opposite in sign

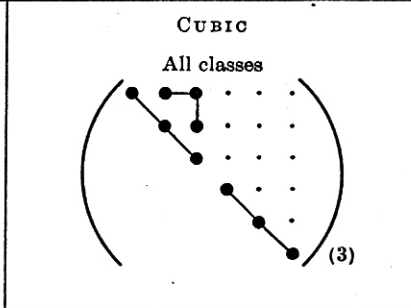
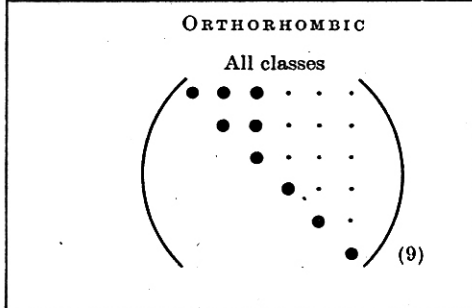
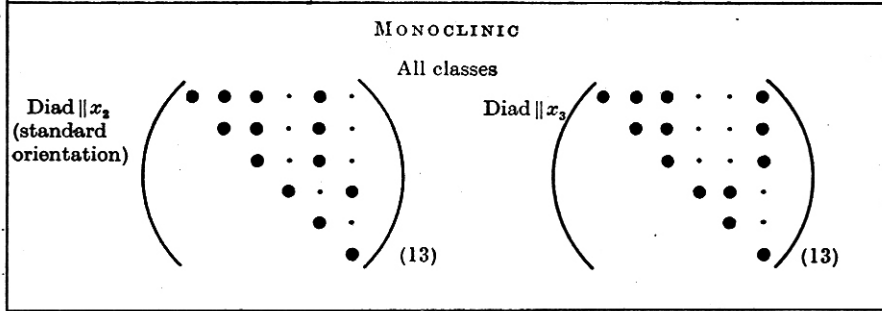
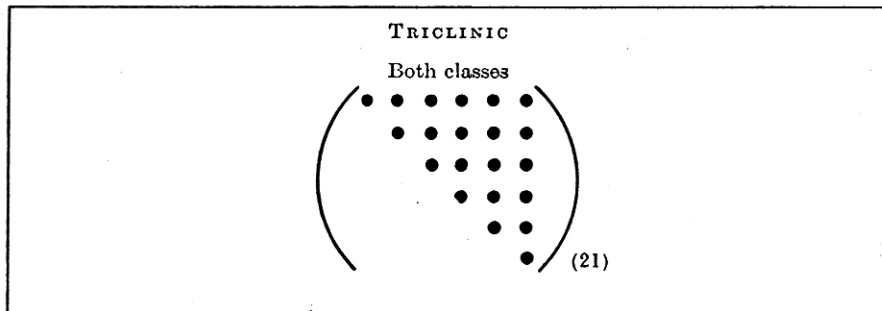
For s ⊙ twice the numerical equal of the heavy dot component to which it is joined

For c ⊙ the numerical equal of the heavy dot component to which it is joined

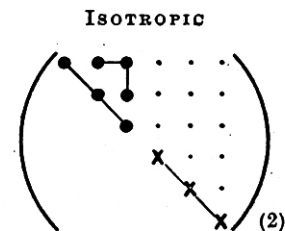
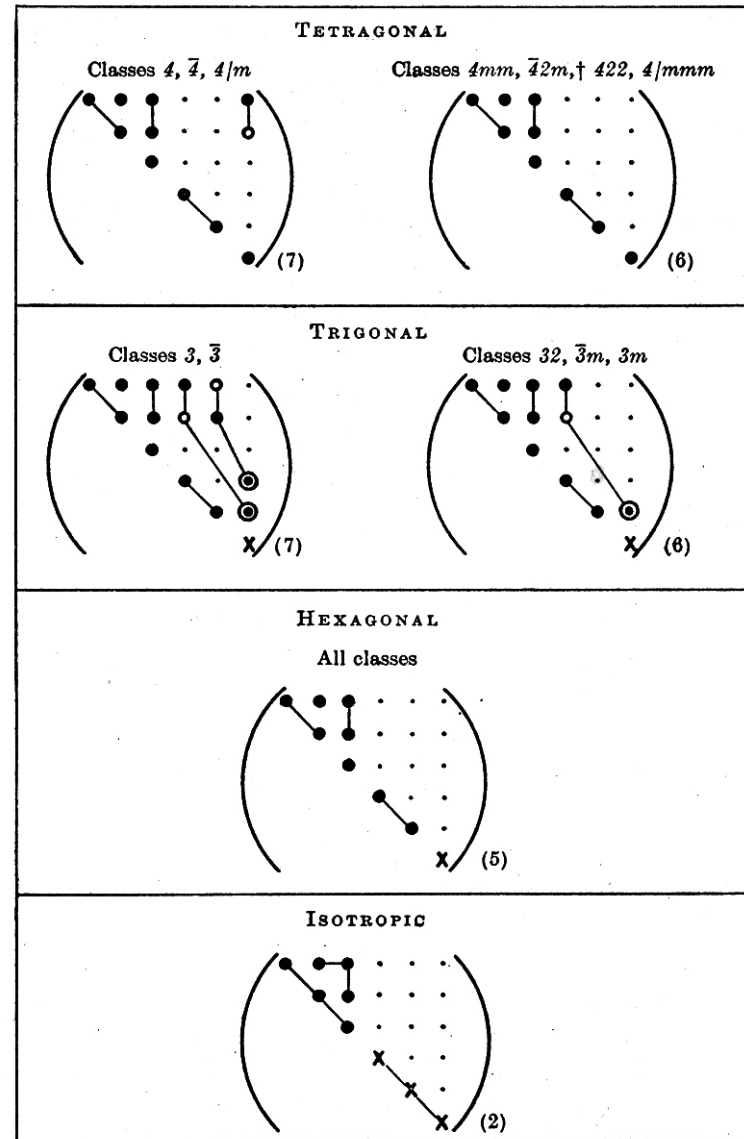
For s X $2(s_{11}-s_{12})$

For c X $\frac{1}{2}(c_{11}-c_{12})$

All the matrices are symmetrical about the leading diagonal.



Effect of Symmetry on the Elasticity Tensors, S, C



General Anisotropic Properties

Objective
Linear
Ferro-
magnets
Non-linear
properties
Electric.
Conduct.
Tensors
Elasticity
Symmetry

- Many different properties of crystals can be described as tensors.
- The rank of each tensor property depends, naturally, on the nature of the quantities related by the property.

Examples of Materials Properties as Tensors

Objective
Linear
Ferro-
magnets
Non-linear
properties
Electric.
Conduct.
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Symmetry

- Table 1 shows a series of tensors that are of importance for material science. The tensors are grouped by rank, and are also labeled (in the last column) by E (equilibrium property) or T (transport property). The number following this letter indicates the maximum number of independent, nonzero elements in the tensor, taking into account symmetries imposed by thermodynamics.
- The Field and Response columns contain the following symbols: $\Delta T =$ *temperature difference*, $\Delta S =$ *entropy change*, $E_i =$ electric field components, $H_i =$ magnetic field components, $\varepsilon_{ij} =$ mechanical strain, $D_i =$ electric displacement, $B_i =$ magnetic induction, $\sigma_{ij} =$ mechanical stress, $\Delta\beta_{ij} =$ change of the impermeability tensor, $j_i =$ electrical current density, $\nabla_j T =$ temperature gradient, $h_i =$ heat flux, $\nabla_j c =$ concentration gradient, $m_i =$ mass flux, $\rho^a_i =$ anti-symmetric part of resistivity tensor, $\rho^s_i =$ symmetric part of resistivity tensor, $\Delta\rho_{ij} =$ change in the component ij of the resistivity tensor, $l_i =$ direction cosines of wave direction in crystal, $G =$ gyration constant,

Property	Symbol	Field	Response	Type#
Tensors of Rank 0 (Scalars)				
Specific Heat	C	ΔT	$T \Delta S$	E1
Tensors of Rank 1 (Vectors)				
Electrocaloric	p_i	E_i	ΔS	E3
Magnetocaloric	q_i	H_i	ΔS	E3
Pyroelectric	p'_i	ΔT	D_i	E3
Pyromagnetic	q'_i	ΔT	B_i	E3
Tensors of Rank 2				
Thermal expansion	α_{ij}	ΔT	ϵ_{ij}	E6
Piezocaloric effect	α'_{ij}	σ_{ij}	ΔS	E6
Dielectric permittivity	κ_{ij}	E_j	D_i	E6
Magnetic permeability	μ_{ij}	H_j	B_i	E6
Optical activity	g_{ij}	$l_i l_j$	G	E6
Magnetoelectric polarization	λ_{ij}	H_j	D_i	E9
Converse magnetoelectric polarization	λ'_{ij}	E_j	B_i	E9
Electrical conductivity (resistivity)	σ_{ij} (ρ_{ij})	E_j (j_j)	j_i (E_i)	T6
Thermal conductivity	K_{ij}	$\nabla_j T$	h_i	T6
Diffusivity	D_{ij}	$\nabla_j c$	m_i	T6
Thermoelectric power	Σ_{ij}	$\nabla_j T$	E_i	T9
Hall effect	R_{ij}	B_j	ρ_i^a	T9

Tensors of Rank 3				
Piezoelectricity	d_{ijk}	σ_{jk}	D_i	E18
Converse piezoelectricity	d'_{ijk}	E_k	ϵ_{ij}	E18
Piezomagnetism	Q_{ijk}	σ_{jk}	B_i	E18
Converse piezomagnetism	Q'_{ijk}	H_k	ϵ_{ij}	E18
Electro-optic effect	r_{ijk}	E_k	$\Delta\beta_{ij}$	E18
Nernst tensor	Σ_{ijk}	$\nabla_j T B_k$	E_i	T27
Tensors of Rank 4				
Elasticity	$s_{ijkl} (c_{ijkl})$	$\sigma_{kl} (\epsilon_{kl})$	$\epsilon_{ij} (\sigma_{ij})$	E21
Electrostriction	γ_{ijkl}	$E_k E_l$	ϵ_{ij}	E36
Photoelasticity	q_{ijkl}	σ_{kl}	$\Delta\beta_{ij}$	E36
Kerr effect	p_{ijkl}	$E_k E_l$	$\Delta\beta_{ij}$	E36
Magnetoresistance	ξ_{ijkl}	$B_k B_l$	ρ_{ij}^s	T36
Piezoresistance	Π_{ijkl}	σ_{kl}	$\Delta\rho_{ij}$	T36
Magnetothermoelectric power	Σ_{ijkl}	$\nabla_j T B_k B_l$	E_i	T54
Second order Hall effect	ρ_{ijkl}	$B_j B_k B_l$	ρ_i^2	T30
Tensors of Rank 6				
Third order elasticity	c_{ijklmn}	$\epsilon_{kl}\epsilon_{mn}$	σ_{ij}	E56

Courtesy of Prof. M. De Graef

$$\begin{pmatrix} \Delta S \\ D_x \\ D_y \\ D_z \\ B_x \\ B_y \\ B_z \\ \epsilon_{xx} \\ \epsilon_{yy} \\ \epsilon_{zz} \\ \epsilon_{yz} \\ \epsilon_{xz} \\ \epsilon_{xy} \end{pmatrix} = \begin{pmatrix} \frac{C}{T} & p_x & p_y & p_z & q_x & q_y & q_z & \alpha'_{xx} & \alpha'_{yy} & \alpha'_{zz} & \alpha'_{yz} & \alpha'_{xz} & \alpha'_{xy} \\ p'_x & \kappa_{xx} & \kappa_{xy} & \kappa_{xz} & \lambda_{xx} & \lambda_{xy} & \lambda_{xz} & d_{xxx} & d_{xyy} & d_{xzz} & d_{xyz} & d_{xxz} & d_{xxy} \\ p'_y & \kappa_{yx} & \kappa_{yy} & \kappa_{yz} & \lambda_{yx} & \lambda_{yy} & \lambda_{yz} & d_{yxx} & d_{yyy} & d_{yzz} & d_{yyz} & d_{yxz} & d_{yxy} \\ p'_z & \kappa_{zx} & \kappa_{zy} & \kappa_{zz} & \lambda_{zx} & \lambda_{zy} & \lambda_{zz} & d_{zxx} & d_{zyy} & d_{zzz} & d_{zuz} & d_{zxx} & d_{zxy} \\ q'_x & \lambda'_{xx} & \lambda'_{xy} & \lambda'_{xz} & \mu_{xx} & \mu_{xy} & \mu_{xz} & Q_{xxx} & Q_{xyy} & Q_{xzz} & Q_{xyz} & Q_{xxz} & Q_{xxy} \\ q'_y & \lambda'_{yx} & \lambda'_{yy} & \lambda'_{yz} & \mu_{yx} & \mu_{yy} & \mu_{yz} & Q_{yxx} & Q_{yyy} & Q_{yzz} & Q_{yyz} & Q_{yxz} & Q_{yxy} \\ q'_z & \lambda'_{zx} & \lambda'_{zy} & \lambda'_{zz} & \mu_{zx} & \mu_{zy} & \mu_{zz} & Q_{zxx} & Q_{zyy} & Q_{zzz} & Q_{zyz} & Q_{zxx} & Q_{zxy} \\ \alpha_{xx} & d'_{xxx} & d'_{xxy} & d'_{xxz} & Q'_{xxx} & Q'_{xxy} & Q'_{xxz} & s_{xxxx} & s_{txyy} & s_{txzz} & s_{txyz} & s_{txxz} & s_{txxy} \\ \alpha_{yy} & d'_{yyx} & d'_{yyy} & d'_{yyz} & Q'_{yyx} & Q'_{yyy} & Q'_{yyz} & s_{yyxx} & s_{yyyy} & s_{yyzz} & s_{yyyz} & s_{yyxz} & s_{yyxy} \\ \alpha_{zz} & d'_{zzx} & d'_{zzy} & d'_{zzz} & Q'_{zzx} & Q'_{zzy} & Q'_{zzz} & s_{zzxx} & s_{zzyy} & s_{zzzz} & s_{zzyz} & s_{zzxz} & s_{zzxy} \\ \alpha_{yz} & d'_{yzx} & d'_{yzy} & d'_{yzz} & Q'_{yzx} & Q'_{yzy} & Q'_{yzz} & s_{yzxx} & s_{yzyy} & s_{yzzz} & s_{yzyz} & s_{yzzx} & s_{yzyx} \\ \alpha_{xz} & d'_{xzx} & d'_{xzy} & d'_{xzz} & Q'_{xzx} & Q'_{xzy} & Q'_{xzz} & s_{xzxz} & s_{xzyy} & s_{xzzz} & s_{xzyz} & s_{xzzz} & s_{xzxy} \\ \alpha_{xy} & d'_{xyx} & d'_{xyy} & d'_{xyz} & Q'_{xyx} & Q'_{xyy} & Q'_{xyz} & s_{xyxx} & s_{xyyy} & s_{xyzz} & s_{xyyz} & s_{xyxz} & s_{xyxy} \end{pmatrix} \begin{pmatrix} \Delta T \\ E_x \\ E_y \\ E_z \\ H_x \\ H_y \\ H_z \\ \sigma_{xx} \\ \sigma_{yy} \\ \sigma_{zz} \\ \sigma_{yz} \\ \sigma_{xz} \\ \sigma_{xy} \end{pmatrix}$$

magnets
 Non-linear properties
 Electric Conduct.
 Tensors
 Elasticity
 Symmetry

- Principal Effects
- Electrocaloric = pyroelectric
- Magnetocaloric = pyromagnetic
- Thermal expansion = piezocaloric
- Magnetoelectric and converse magnetoelectric
- Piezoelectric and converse piezoelectric
- Piezomagnetic and converse piezomagnetic

Courtesy of Prof. M. De Graef

$$\Delta S = \frac{C}{T} \Delta T + p_i E_i + q_i H_i + \alpha'_{ij} \sigma_{ij};$$

$$D_i = p'_i \Delta T + \kappa_{ij} E_j + \lambda_{ij} H_j + d_{ijk} \sigma_{jk};$$

$$B_i = q'_i \Delta T + \lambda'_{ij} E_j + \mu_{ij} H_j + Q_{ijk} \sigma_{jk};$$

$$\epsilon_{ij} = \alpha_{ij} \Delta T + d'_{ijk} E_k + Q'_{ijk} H_k + s_{ijkl} \sigma_{kl}.$$

Non-linear
properties

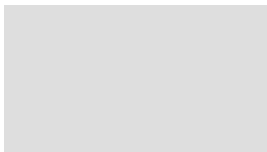
Electric.
Conduct.

Tensors

Elasticity
Symmetry



Principal Effects



1st rank cross effects



2nd rank cross effects



3rd rank cross effects

Courtesy of Prof. M. De Graef

$$\begin{pmatrix} \Delta S \\ D_x \\ D_y \\ D_z \\ B_x \\ B_y \\ B_z \\ \epsilon_{xx} \\ \epsilon_{yy} \\ \epsilon_{zz} \\ \epsilon_{yz} \\ \epsilon_{xz} \\ \epsilon_{xy} \end{pmatrix} = \begin{pmatrix} \frac{C}{T} & p_x & p_y & p_z & q_x & q_y & q_z & \alpha'_{xx} & \alpha'_{yy} & \alpha'_{zz} & \alpha'_{yz} & \alpha'_{xz} & \alpha'_{xy} \\ p'_x & \kappa_{xx} & \kappa_{xy} & \kappa_{xz} & \lambda_{xx} & \lambda_{xy} & \lambda_{xz} & d_{xxx} & d_{xyy} & d_{xzz} & d_{xyz} & d_{xxz} & d_{xxy} \\ p'_y & \kappa_{yx} & \kappa_{yy} & \kappa_{yz} & \lambda_{yx} & \lambda_{yy} & \lambda_{yz} & d_{yxx} & d_{yyy} & d_{yzz} & d_{yyz} & d_{yxz} & d_{yyx} \\ p'_z & \kappa_{zx} & \kappa_{zy} & \kappa_{zz} & \lambda_{zx} & \lambda_{zy} & \lambda_{zz} & d_{zxx} & d_{zyy} & d_{zzz} & d_{zyz} & d_{zxx} & d_{zxy} \\ q'_x & \lambda'_{xx} & \lambda'_{xy} & \lambda'_{xz} & \mu_{xx} & \mu_{xy} & \mu_{xz} & Q_{xxx} & Q_{xyy} & Q_{xzz} & Q_{xyz} & Q_{xxz} & Q_{xxy} \\ q'_y & \lambda'_{yx} & \lambda'_{yy} & \lambda'_{yz} & \mu_{yx} & \mu_{yy} & \mu_{yz} & Q_{yxx} & Q_{yyy} & Q_{yzz} & Q_{yyz} & Q_{yxz} & Q_{yyx} \\ q'_z & \lambda'_{zx} & \lambda'_{zy} & \lambda'_{zz} & \mu_{zx} & \mu_{zy} & \mu_{zz} & Q_{zxx} & Q_{zyy} & Q_{zzz} & Q_{zyz} & Q_{zxx} & Q_{zxy} \\ \alpha_{xx} & d'_{xxx} & d'_{xxy} & d'_{xxz} & Q'_{xxx} & Q'_{xxy} & Q'_{xxz} & s_{xxxx} & s_{xxyy} & s_{xxzz} & s_{txyz} & s_{txxz} & s_{txxy} \\ \alpha_{yy} & d'_{yyx} & d'_{yyy} & d'_{yyz} & Q'_{yyx} & Q'_{yyy} & Q'_{yyz} & s_{yyxx} & s_{yyyy} & s_{yyzz} & s_{yyyz} & s_{yyxz} & s_{yyxy} \\ \alpha_{zz} & d'_{zzx} & d'_{zzy} & d'_{zzz} & Q'_{zzx} & Q'_{zzy} & Q'_{zzz} & s_{zzxx} & s_{zzyy} & s_{zzzz} & s_{zzyz} & s_{zzxz} & s_{zzxy} \\ \alpha_{yz} & d'_{yzx} & d'_{yzy} & d'_{yzz} & Q'_{yzx} & Q'_{yzy} & Q'_{yzz} & s_{yzxx} & s_{yzyy} & s_{yzzz} & s_{yzyz} & s_{yzzx} & s_{yzyx} \\ \alpha_{xz} & d'_{xzx} & d'_{xzy} & d'_{xzz} & Q'_{xzx} & Q'_{xzy} & Q'_{xzz} & s_{xzxz} & s_{xzyy} & s_{xzzz} & s_{xzyz} & s_{xzzz} & s_{xzxy} \\ \alpha_{xy} & d'_{xyx} & d'_{xyy} & d'_{xyz} & Q'_{xyx} & Q'_{xyy} & Q'_{xyz} & s_{xyxx} & s_{xyyy} & s_{xyzz} & s_{xyyz} & s_{xyxz} & s_{xyxy} \end{pmatrix} \begin{pmatrix} \Delta T \\ E_x \\ E_y \\ E_z \\ H_x \\ H_y \\ H_z \\ \sigma_{xx} \\ \sigma_{yy} \\ \sigma_{zz} \\ \sigma_{yz} \\ \sigma_{xz} \\ \sigma_{xy} \end{pmatrix}$$

magnets
 Non-linear
 properties
 Electric.
 Conduct.
 Tensors
 Elasticity
 Symmetry

General crystal symmetry shown above.

Courtesy of Prof. M. De Graef

$$\begin{pmatrix} \Delta S \\ D_x \\ D_y \\ D_z \\ B_x \\ B_y \\ B_z \\ \epsilon_{xx} \\ \epsilon_{yy} \\ \epsilon_{zz} \\ \epsilon_{yz} \\ \epsilon_{xz} \\ \epsilon_{xy} \end{pmatrix} = \begin{pmatrix} C \\ T & p_x & p_x & p_z & q_x & q_x & q_z & \alpha_{xx} & \alpha_{xx} & \alpha_{zz} & 0 & 0 & 0 \\ p_x & \kappa_{xx} & 0 & 0 & \lambda_{xx} & \lambda_{xy} & 0 & 0 & 0 & 0 & d_{xyz} & d_{xxz} & 0 \\ p_x & 0 & \kappa_{xx} & 0 & -\lambda_{xy} & \lambda_{xx} & 0 & 0 & 0 & 0 & -d_{xxz} & d_{xyz} & 0 \\ p_z & 0 & 0 & \kappa_{zz} & 0 & 0 & \lambda_{zz} & d_{zxx} & d_{zxx} & d_{zzz} & 0 & 0 & 0 \\ q_x & \lambda_{xx} & -\lambda_{xy} & 0 & \mu_{xx} & 0 & 0 & 0 & 0 & 0 & Q_{xyz} & Q_{xxz} & 0 \\ q_x & \lambda_{xy} & \lambda_{xx} & 0 & 0 & \mu_{xx} & 0 & 0 & 0 & 0 & -Q_{xxz} & Q_{xyz} & 0 \\ q_z & 0 & 0 & \lambda_{zz} & 0 & 0 & \mu_{zz} & Q_{zxx} & Q_{zxx} & Q_{zzz} & 0 & 0 & 0 \\ \alpha_{xx} & 0 & 0 & d_{zxx} & 0 & 0 & Q_{zxx} & s_{xxxx} & s_{txyy} & s_{txzz} & 0 & 0 & s_{txxy} \\ \alpha_{xx} & 0 & 0 & d_{zxx} & 0 & 0 & Q_{zyy} & s_{txyy} & s_{xxxx} & s_{txzz} & 0 & 0 & -s_{txxy} \\ \alpha_{zz} & 0 & 0 & d_{zzz} & 0 & 0 & Q_{zzz} & s_{txzz} & s_{txzz} & s_{zzzz} & 0 & 0 & 0 \\ 0 & d_{xyz} & -d_{xxz} & 0 & Q_{xyz} & -Q_{xxz} & 0 & 0 & 0 & 0 & s_{txxz} & -s_{txyz} & 0 \\ 0 & d_{xxz} & d_{xyz} & 0 & Q_{xxz} & Q_{xyz} & 0 & 0 & 0 & 0 & s_{txyz} & s_{txxz} & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 & s_{txxy} & -s_{txxy} & 0 & 0 & 0 & s_{txxy} \end{pmatrix} \begin{pmatrix} \Delta T \\ E_x \\ E_y \\ E_z \\ H_x \\ H_y \\ H_z \\ \sigma_{xx} \\ \sigma_{yy} \\ \sigma_{zz} \\ \sigma_{yz} \\ \sigma_{xz} \\ \sigma_{xy} \end{pmatrix}$$

Non-linear
properties

Electric.
Conduct.

Tensors

Elasticity

Symmetry

Point group 4

Courtesy of Prof. M. De Graef

$$\begin{pmatrix} \Delta S \\ D_x \\ D_y \\ D_z \\ B_x \\ B_y \\ B_z \\ \epsilon_{xx} \\ \epsilon_{yy} \\ \epsilon_{zz} \\ \epsilon_{yz} \\ \epsilon_{xz} \\ \epsilon_{xy} \end{pmatrix} = \begin{pmatrix} \frac{C}{T} & 0 & 0 & 0 & 0 & 0 & 0 & \alpha & \alpha & \alpha & 0 & 0 & 0 \\ 0 & \kappa & 0 & 0 & \lambda & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & \kappa & 0 & 0 & \lambda & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & \kappa & 0 & 0 & \lambda & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & \lambda & 0 & 0 & \mu & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & \lambda & 0 & 0 & \mu & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & \lambda & 0 & 0 & \mu & 0 & 0 & 0 & 0 & 0 & 0 \\ \alpha & 0 & 0 & 0 & 0 & 0 & 0 & s_{xxxx} & s_{xxyy} & s_{xxyy} & 0 & 0 & 0 \\ \alpha & 0 & 0 & 0 & 0 & 0 & 0 & s_{xxyy} & s_{xxxx} & s_{xxyy} & 0 & 0 & 0 \\ \alpha & 0 & 0 & 0 & 0 & 0 & 0 & s_{xxyy} & s_{xxyy} & s_{xxxx} & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & s_{yzyz} & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & s_{yzyz} & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & s_{yzyz} \end{pmatrix} \begin{pmatrix} \Delta T \\ E_x \\ E_y \\ E_z \\ H_x \\ H_y \\ H_z \\ \sigma_{xx} \\ \sigma_{yy} \\ \sigma_{zz} \\ \sigma_{yz} \\ \sigma_{xz} \\ \sigma_{xy} \end{pmatrix}$$

Conduct.

Tensors

Elasticity

Symmetry

Point group $m\bar{3}m$

Note how many fewer independent coefficients there are!
 Note how the center of symmetry eliminates many of the properties, such as pyroelectricity

Courtesy of Prof. M. De Graef

Homogeneity

Objective
Linear
Ferro-
magnets
Non-linear
properties
Electric.
Conduct.
Tensors
Elasticity
Symmetry

- Stimuli and responses of interest are, in general, not scalar quantities but tensors. Furthermore, some of the properties of interest, such as the plastic properties of a material, are far from linear at the scale of a polycrystal. Nonetheless, they can be treated as linear at a suitably local scale and then an averaging technique can be used to obtain the response of the polycrystal. The local or microscopic response is generally well understood but the validity of the averaging techniques is still controversial in many cases. Also, we will only discuss cases where a homogeneous response can be reasonably expected.
- There are many problems in which a non-homogeneous response to a homogeneous stimulus is of critical importance. Stress-corrosion cracking, for example, is a wildly non-linear, non-homogeneous response to an approximately uniform stimulus which depends on the mechanical and electro-chemical properties of the material.