A Very Abbreviated Introduction to Powder Diffraction

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Outline

- Stuff you should know:
  - Diffraction from single crystals
  - X-rays vs Neutrons; Types of scattering
- Diffraction from powders
- Why do we use powder diffraction?
- Instruments for powder diffraction collection
- Materials effects in powder diffraction
- Crystallographic analysis of powder diffraction data
- Appendices:
  - More on peak shapes
  - More on crystallography
  - Where to go for more information
Single Crystal Diffraction Review
Single crystal diffraction: the movie

Video from Diamond synchrotron (U.K.)
Bragg scattering in single crystals

- Reflection positions (Bragg’s law)
  \[ 2d \sin \theta = n \lambda \]

Note that \( d \) above is a reciprocal space quantity (actually \( d = 1/d^* \)), \( |d^*| = |ha^* + kb^* + lc^*| \),
better: \( \lambda = 4 \pi \sin \theta / Q \)

- Reflection intensities are related to the square of “structure factor”, \( F_{hk\ell} \)
  \[
  F_{hk\ell} \propto \sum f_j \exp[2\pi i(hx_j + ky_j + lz_j)] \exp(-U_j Q^2/2) \]
  sum over all atoms in the crystal (or unit cell).
  \( x_j, y_j, z_j \): position of atom in unit cell
  \( U_j \): atom displacements
  \( f_j \) (or \( b_j \)): scattering power of atom
Coherent Atomic Scattering Power (diffraction)

**X-rays**: The scattering power (form factor, $f_i$) of an atom depends on the number of electrons in the atom and $Q$ ($Q \propto \sin \theta / \lambda$)
- X-ray scattering changes near absorption edges

**Neutrons**: The scattering power (scattering length, $b_i$) of an atom depends on the isotope and is independent of $Q$
- A few isotopes scatter with opposite phase to most, for these we write $b_i$ (f) as negative
- Some isotopes atoms have neutron resonances (similar to x-rays)
- Magnetic scattering is from electrons; $f_M(Q)$ similar to x-rays

**Structure factors**: $F_{hkl} = n \sum f_i \exp[2\pi i(hx_i + ky_i + lz_i)] \exp(-U_i Q^2/2)$
Comparison of Neutron and X-ray Atomic Scattering Powers
Neutrons and x-rays “see” atoms differently
Resonant scattering: scattering at a resonance edge causes atoms to “light up”

Experiments are sometimes performed at wavelengths close to absorption edges to enhance the scattering from particular elements

**X-rays**

The x-ray form factor has in fact three components:

- \( f(Q) + f'(\lambda) + i f''(\lambda) \)
  - \( f \) is determined by \( Q \) and the number of electrons in an atom and is independent of wavelength
  - \( f' \) and \( f'' \) are small except at wavelengths very close to an atom’s absorption edge

At wavelengths close to an edge absorption becomes high; fluorescence occurs above the edge.

**Neutrons**

Scattering lengths for most atoms are wavelength-independent.

A similar “resonant scattering” type experiment can sometimes be performed comparing samples containing different isotopes ($\$$ to $\$$\$$)$

A few isotopes (mostly lanthanides and actinides) have adsorption edges at accessible wavelengths.

- This is usually a curse rather than a blessing: it makes TOF neutron scattering had to analyze
Types of scattering
Coherent vs Incoherent Scattering

- Coherent scattering -- since phase is retained, scattering occurs with interference between nearby atoms
  - atom-atom correlation info obtained: diffraction

- Incoherent scattering -- probe phase is randomized, no interference
  - self-correlation (simpler analysis): spectroscopy

Elastic vs Inelastic Scattering

- Elastic scattering -- No transfer of energy between atoms and probe, only momentum

- Inelastic scattering -- probe loses energy to atoms (for neutrons also possible to gain energy)
## Types of neutron scattering measurements

<table>
<thead>
<tr>
<th></th>
<th>Elastic Scattering</th>
<th>Inelastic Scattering</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Coherent Scattering</strong></td>
<td>Diffraction (structural studies)</td>
<td>Phonons, magnons… (collective excitations)</td>
</tr>
<tr>
<td><strong>Incoherent Scattering</strong></td>
<td>QENS (diffusion, low barrier motion)</td>
<td>Neutron Spectroscopies (atomic vibrations)</td>
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</tbody>
</table>
Incoherent and inelastic scattering create background. *This is usually significant only with neutrons and most commonly for powder (less so single crystal) diffraction.*

With neutrons, some type of atoms have large incoherent cross sections (phase is lost during the scattering).

Hydrogen (not deuterium) is the poster child for this: it has a huge incoherent scattering cross-section (~80 barns) that tends to overpower coherent scattering (typically <1 barns for most atoms). Samples with more than a few atom % (not mass %!) will have a significant background.
Powder Diffraction
Diffraction from random polycrystalline material

In a sufficiently large, randomly oriented polycrystalline sample (e.g. a powder) contains a very large number of crystallites.

A beam impinging on the sample will find a representative number of crystallites in the right orientation for diffraction.

Diffraction occurs only at specific angles, those where Bragg’s Law is satisfied.
Bragg cones in powder diffraction

Since there is a random distribution of crystals then diffraction occurs in all directions for each Bragg angle, $2\theta$, thus powder diffraction arises in cones.

All reflections occurring at a single $2\theta$ value (as well as reflections at nearly the same value) are superimposed and only the sum of their intensities can be measured. For this reason a powder diffraction pattern gives less information than a single crystal measurement.
Why do we do powder diffraction?

- Learn where the atoms are (single crystals, when available and appropriate, are better for this.)

- Determine the chemical phase(s) in a sample

- Measure lattice constants

- Quantify the components of a mixture

- Learn about physical specimen characteristics such as stress, preferred orientation or crystallite sizes

- Occupancies of elements amongst crystallographic sites (usually needs neutrons)
Types of Powder Diffraction Measurements
Measuring powder diffraction

- Angular dispersion: a single detector is moved over a range of $2\theta$ angles.
  - Sample irradiated with monochromatic radiation
With an area detector, a complete powder diffraction pattern can be collected in a fraction of a second.
- Fast
- Medium resolution
- High background
Highest resolution requires high collimation. Optimal is a crystal analyzer between the sample and detector: 11-BM Diffractometer

Huber 480 rotation stage:
- high precision (~0.35arcsec)
- high accuracy (~1arcsec)
- slew or step scans

12 analyzer array
- Si(111) crystals
- LaCl$_3$ scintillator detectors
- 2° apart in 2Θ.

Complete pattern is measured in <1 hour
11-BM Robotic Data Collection

(Matt Suchomel: [http://youtu.be/sowojskY7c4](http://youtu.be/sowojskY7c4) or search APS 11-BM on YouTube)
Reactor Source Neutron Diffraction

Producing neutrons by fission

Slow Neutron → 235U nucleus

Fission Fragments → Fast Neutrons are slowed by collisions in the moderator (C, H2O, D2O)

Steady State Technique

Sample

Monochromator Crystal

I(λ)

Small Δλ used

Source on all the time

I(t) on Sample

I(t) at Detector

Time →
Powder Instruments: Constant Wavelength

beamline HB2a at HFIR

beamline BT1 at NIST (NCNR)
Comparison of $\text{Tb}_2\text{TiO}_7$ with x-rays & neutrons

Intensity

$2\Theta @ 1.2 \text{ Å}$

X-ray

Neutron
Spallation Sources

A pulse of protons impacting the target produces a shower of fast neutrons that are slowed down in a moderator. A new pulse is created ~30 times/sec

**Producing neutrons by spallation**

- Fast proton
- Heavy nucleus (Ta, U, Hg)
- Fast Neutrons are slowed by collisions in a moderator (CH₄, H₂O, D₂O)

Each pulse of neutrons contains a broad spectrum of neutron energies.
**Time of Flight Diffraction**

Time of flight diffraction uses the fact that neutrons with different energies (velocities) have different wavelengths, $\lambda = \frac{h}{mv}$ (de Broglies relationship).
Time-of-flight \((2\sin \theta = \lambda)\)

\(\lambda \text{ varies}\)

\(2\theta \text{ fixed}\)

(Pulsed sources: e.g. SNS)
Neutron Powder Diffraction with Spallation Source

- Spallation source provides a broad band of wavelengths in sharp pulses
  - TOF detection allows measurement of intensity versus wavelength
  - Each detector provides a full diffraction pattern
  - Data collection times:
    • Seconds to hours

NPDF instrument at LANSCE (Los Alamos)
3rd Generation TOF Instruments: High Intensity and High Resolution

- **GEM (ISIS TS-I)**
- **WISH**
  A high-resolution magnetic diffractometer for TS-II
- **Super HRPD (JSNS)**
Understanding Materials Effects in Powder Diffraction
Materials effects on Powder Diffraction

Peak broadening:

- Crystallite size:
  - What happens when crystals become small?

- Residual Stress (Strain)
  - What happens if matrix effects do not allow crystallites to equilibrate lattice parameters?

Note that these effects can vary in different crystallographic directions: anisotropic peak broadening (best handled in GSAS-II)
Crystallite Size Broadening

The Fourier transform (FT) from an infinite array of regularly spaced objects is an array of delta functions.

The FT from a finite length array is broadened.

The finite sizes of crystallites will broaden all orders of reflections equally in units of Q (\( \propto d^* \))

- differing reciprocal space directions may have differing amounts of broadening, if crystallites dimensions are not isotropic on average

Crystallite Size Broadening can produce Lorentzian peak shapes (common) or Gaussian peak shapes (uncommon) or a combination of both.
Crystallite Size Broadening

\[ \Delta d^* = \text{constant} \]

\[ \Delta d^* = \frac{\Delta d}{d^2} = \frac{\Delta \Theta \cot \Theta}{d} \]

\[ \Delta 2\Theta = \frac{\lambda \Delta d}{d^2 \cos \Theta} \]

GSAS fits crystallite broadening with two profile terms:
- LX \(\rightarrow\) Lorentzian
- GP \(\rightarrow\) Gaussian

Relation between avg. size (\(p\)) and GSAS terms:

\[ p = \frac{18000 K \lambda}{\pi LX} \quad \text{and} \quad p = \frac{18000 K \lambda}{\pi \sqrt{GP}} \]

\(K \approx 1\) (Scherrer constant, related to crystal shape)

Microstrain Broadening

When a material has residual stresses present, some crystallites are compressed. This must be balanced by other crystallites that are stretched (because $\sum F = ma = 0$)

This leads to a range of lattice constants.

The spread between diffraction locations for the maximum and minimum lattice constant increases linearly with Q ($\Delta Q/Q$ or $\Delta d/d = \text{constant}$)
Microstrain Broadening

\[
\frac{\Delta d}{d} = \text{constant}
\]

\[
\frac{\Delta d}{d} = \frac{\Delta d^*}{d^*} = \Delta \Theta \cot \Theta
\]

\[
\Delta 2\Theta = \frac{2\Delta d}{d} \tan \Theta
\]

GSAS fits strain broadening with two profile terms:
- LY -> Lorentzian (most common)
- GU -> Gaussian

(note that GU also has an instrumental contribution)


Relation between strain (as percentage) and GSAS terms:

\[
S = 100\% \frac{\pi}{18000} \text{LY}
\]

\[
S = 100\% \frac{\pi}{18000} \sqrt{GU - GU_i}
\]

where \(GU_i\) accounts for the instrumental contribution
Fitting of Powder Diffraction Data (Rietveld Analysis)
Why did Crystallography Revolutionize Science?

1. Crystallography was the first scientific technique that provided direct information about molecular structure
   - Early work was intuitive: structures assigned based on patterns and symmetry (some results predate X-rays!)

2. X-ray and neutron diffraction observations can be modeled very accurately directly when the molecular structure is known

3. Diffraction can provide a very large number of independent observations
   - Probability of finding an incorrect structure model that is both plausible and in good agreement with the diffraction observations is very small (but not zero!)

4. Computer-assisted least-squares optimization allows structural models to be improved, limited only by the quality of the data

5. Statistical and brute-force techniques overcome the incomplete nature of diffraction observations (direct methods vs. “the phase problem”).

100+ years later, no other technique offers as much power for learning about molecular structure!
We perform an experiment:
- Get lots of intensity and position measurements in a diffraction measurement: what do they tell us?

Obtain an unit cell that fits the diffraction positions (indexing)

“Solve the structure”: determine an approximate model to match the intensities

Add/modify the structure for completeness & chemical sense

Optimize the structure (model) to obtain the best fit to the observed data
- This is usually done with Gauss-Newton least-squares fitting
- Parameters to be fit are structural and may account for other experimental effects

Least Squares gives us a Hessian matrix; inverse is variance-covariance matrix which gives uncertainties in the parameters
Crystallography from powder diffraction: before Rietveld

How did crystallographers use powder diffraction data?

- Avoided powder diffraction
- Manually integrate intensities
  - discard peaks with overlapped reflections
  - rewrote single-crystal software to refine using sums of overlapped reflections

Simulation of powder diffraction data was commonly done

- Qualitative reasoning: similarities in patterns implied similar structures
- Visual comparison between computed and observed structure verifies approximate model
- Fits, where accurate (& precise) models were rarely obtained

Error propagation was difficult to do correctly (but not impossible)
Hugo Rietveld’s technique

- Hugo Rietveld realized that if a pattern could be modeled, the fit between a computed pattern and observed data could be optimized.

  - Similar to single-crystal diffraction, except that now “experiment dependent parameters” must now be fit as well.
    - Background
    - Peak broadening
    - Lattice constants
  - Must have approximate model to start
  - Fewer data are available (usually)
Calculation of Powder Diffraction: Graphical Example

1) Generate reflection list
2) Compute $F_{hkl}$ from model
3) Peak heights are generated from $|F_{hkl}|^2 \times$ multiplicity
4) Convolute peaks & add background
5) Optimize model, peak widths, etc. to improve fit

<table>
<thead>
<tr>
<th>$hkl$</th>
<th>mult</th>
<th>D-space</th>
<th>$F_{hkl}$</th>
<th>phase</th>
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<td>0.29</td>
<td>0</td>
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<td>180</td>
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<td>1.417</td>
<td>1.827</td>
<td>180</td>
</tr>
</tbody>
</table>
Hugo Rietveld in the Petten Reactor (~1987)
Single crystal fitting

Minimize equation $\Sigma w_i [y_i - Y(x_i, p)]^2$ where

**Data:** $y_i = F_{hkl} (\text{obs})$

$y_i =$ observed powder diffraction intensities

**Model:** $Y(x_i, p) = F_{hkl} (\text{calc})$

$Y(x_i, p) =$ computed diffraction intensities from ($F_{hkl} (\text{calc})$, background model, profile convolution, preferred orientation correction…

**Parameters** ($\rho_1, \rho_2, \rho_3… \rho_m$):
atomic coordinates, displacement (T) factors

+ lattice parameters
+ “experimental” parameters for peak shapes, background…
Hugo Rietveld’s other breakthrough

- Based on intensities from the model, estimates for $F_{hkl}$ can be made, even when reflections are completely overlapped:

  Location 1:
  - 20% to A
  - 40% to B
  - 40% to C

  Location 2:
  - 100% to C
Rietveld Applications

- Crystallographic structure determination
- Quantify amounts of crystalline phases
  - (Amorphous content possible indirectly)
- Engineering properties
  - Residual stress/Crystallite sizes
  - Preferred orientation
- Lattice constant determination
What sort of data are needed for Rietveld Analysis?

- Must be possible to fit peak shapes
- Q range and resolution demands dictated by structural complexity
- Data from lab instruments should be used *with caution* for structure determination
- Neutron data are usually necessary for occupancy determination
Disadvantage of Rietveld: Many non-structural parameters need to be fit

- **Background**
  - fixed
  - functions
- **Peak shape**
  - "fundamental parameters"
  - functions
- **Lattice constants**
  - zero correction
  - flat plate terms

- **Scaling**
  - Phase fractions
- **Structural parameters**
  - atom positions
  - occupancies
  - displacement parameters
- **Preferential Orientation**
- **Absorption**

*Powder diffraction offers fewer observations and worse peak-to-background than single crystal diffraction*
Limitations of Rietveld

- Rietveld can only discern parameters that have effects on the powder diffraction pattern
  - Some things cannot ever be determined from powder diffraction:
    - Absolute configuration
    - “in symmetry plane” magnetic moment directions

- If two parameters have approximately the same effect on the powder diffraction pattern, they correlate and they cannot be differentiated (e.g. occupancies & displacement parameters)
The physics that determine peak profiles

Common factors
- Instrumental Resolution
- Axial Divergence (Low Angle Asymmetry)
- Sample placement/transparency
- Crystallite Broadening*
- Strain Broadening*

* Note that these effects can vary for different classes of reflections (anisotropic peak broadening)

Less common factors
- Stacking faults*
- Modulated Structures*
- Coherence differing by atom type*
- Compositional inhomogeneity!

* Hard to model
! Sometimes impossible to model

Results of these factors are convoluted to produce the observed peak shape.
Appendix 1: more on peak profiles
Three different approaches to reproducing peak shapes have been used:

- **Empirical functions**
  
  Functions are chosen simply because they are computationally simple and fit peaks well. The parameters cannot be interpreted because they have no relationship to the underlying physics of diffraction.

- **Physically-based parameters**
  
  Functions are based on the physical phenomena. Parameters are usually found empirically, but often have a physical meaning.

- **“Fundamental Parameters”**
  
  Functions and where possible parameter values are determined from diffraction physics. The only adjustable parameters are those related to sample properties.
Lorentz (Cauchy) and Gaussian Broadening Functions

Most instrument & sample broadening contributions are Lorentzian or Gaussian

Normalized Gaussian

\[ G(\Delta T, \Gamma_G) = \frac{4\ln 2}{\pi \Gamma_G^2} \exp \left[ -\frac{4 \ln 2 (\Delta T)^2}{\Gamma_G^2} \right] \]

Normalized Lorentzian

\[ L(\Delta T, \gamma_L) = \frac{2}{\pi \gamma_L} \frac{1}{1 + \left( \frac{2\Delta T}{\gamma_L} \right)^2} \]

Note that peak widths vary so \( \Gamma_G \) and \( \gamma_L \) are both functions of \( Q \)

Gaussian & Lorentzian functions compared. Both curves have same FWHM & area, but note the much longer tails for the Lorentzian.
Voigt vs. Pseudo-Voigt

A Gaussian convoluted with a Lorentzian function is a Voigt function, however the Voigt is slow to compute and the derivatives are messy. Few Rietveld programs implement a Voigt.

The “pseudo-Voigt” is the weighted sum of a Gaussian & Lorentzian function – approximation is normally pretty good

Fractions of each function depend on the relative widths of each [see mixing factor ($\eta$) in GSAS manual, $\eta=0$ is Gaussian, $\eta=1$ is Lorentzian]
CW: Variation of FWHM with Q

**Gaussian**
- U, V & W are Cagliotti terms, derived as instrument response function for CW neutron diffraction. Incomplete for x-rays.
- P is a crystallite broadening contribution

**Lorentzian**
- X is crystallite broadening
- Y is strain broadening

\[ \gamma = \frac{X}{\cos \Theta} + Y \tan \Theta \]

**N.B. no instrumental broadening term**
Axial Divergence (Low Angle Asymmetry)

Work of Finger, Cox & Jephcoat, based on derivations by van Laar & Yelon

Debye-Scherrer cone

2θ Scan

H

Slit

FCJ:
Convolute profile with this curve

2θ_{\text{min}}

2θ_i

2θ_{\text{Bragg}}

Depend on slit & sample “heights” wrt diffr. radius
H/L & S/L - parameters in function
(typically 0.005 - 0.020)
F-C-J: Example

- The Finger-Cox-Jephcoat correctly models the effective shift of the peak due to axial divergence.

Note: the “competition,” the split Pearson VII (empirical), does not model this effect at all!
Sample Displacement & Transparency

In **Bragg-Brentano geometry**, samples are ideally placed exactly at rotation axis and all diffraction occurs from sample surface (highly absorbing sample). Neither is commonly true.

- Peak centers are shifted by
  - Sample Displacement (SHFT), $S_s$
  - Sample transparency (TRNS), $T_s$

$$\Delta T' = \Delta T + S_s \cos \Theta + T_s \sin 2\Theta$$

$$\text{displacement} = -\frac{\pi R S_s}{36000} \quad \mu_{\text{eff}} = -\frac{9000}{\pi R T_s}$$

*R is diffractometer radius*

*These corrections correlate very highly with the zero correction for $2\theta$, ZERO. *Do not refine this too.*

**Parallel-Beam instruments** *(neutron or synchrotron)* are very tolerant of displacement and transparency. Never refine SHFT or TRNS, but do refine ZERO (correction to $2\theta$).
When Strain Differs by Reflection Class: 
“Anisotropic peak broadening”

Strain may be anisotropic

- *think of a layered material where the layers can be pulled apart without much effort, but the layers themselves are quite “hard” (resistant to applied forces).*

- Such a material will be “squishy” in the layer direction and rigid in the other two (more broadening in the squishy direction.)


- Restricts strain components in terms of 1st & 2nd-order terms allowed by lattice symmetry
Anisotropic strain broadening terms

Broadening – as variance

$$\sigma^2(M_{hkl}) = \sum_{HKL} S_{HKL} h^H k^K l^L , \ H + K + L = 4$$

General expression – triclinic – 15 terms

$$\sigma^2(M_{hkl}) = S_{400} h^4 + S_{040} k^4 + S_{004} l^4 + 3\left(S_{220} h^2 k^2 + S_{202} h^2 l^2 + S_{022} k^2 l^2 \right) +$$
$$2\left(S_{310} h^3 k + S_{103} h l^3 + S_{031} k^3 l + S_{130} h k^3 + S_{301} h^3 l + S_{013} k l^3 \right) +$$
$$4\left(S_{211} h^2 k l + S_{121} h k^2 l + S_{112} h k l^2 \right)$$

Symmetry effects – monoclinic (b unique) – 9 terms

$$\sigma^2(M_{hkl}) = S_{400} h^4 + S_{040} k^4 + S_{004} l^4 + 3S_{202} h^2 l^2 + 3\left(S_{220} h^2 k^2 + S_{022} k^2 l^2 \right) +$$
$$2\left(S_{301} h^3 l + S_{103} h k^3 \right) + 4S_{121} h k^2 l$$
Anisotropic strain broadening terms

Symmetry effects – orthorhombic - mmm – 6 terms

\[ \sigma^2(M_{hkl}) = S_{400} h^4 + S_{040} k^4 + S_{004} l^4 + 3\left( S_{220} h^2 k^2 + S_{202} h^2 l^2 + S_{022} k^2 l^2 \right) \]

Tetragonal – 4/mmm – 4 terms

\[ \sigma^2(M_{hkl}) = S_{400} (h^4 + k^4) + S_{004} l^4 + 3S_{220} h^2 k^2 + 3S_{202} (h^2 l^2 + k^2 l^2) \]

Cubic – m3m – 2 terms

\[ \sigma^2(M_{hkl}) = S_{400} (h^4 + k^4 + l^4) + 3S_{220} (h^2 k^2 + h^2 l^2 + k^2 l^2) \]

Trigonal, Rhombohedral & Hexagonal – it depends – 3 or 4 terms
Appendix 2: more on crystallography
The Lattice

- Crystals are constructed from repeated arrangements of atoms.
- Crystalline structure can be described as set of “identical boxes” stacked in 3D; the contents of each box is identical (exception: quasicrystals)
  - **A lattice** is a mathematical concept where each lattice point describes an identical environment; lattice points are the corners of the “identical boxes.”

Commonly used phrases such as “lattice compound” or “interstitials in the lattice” misuse the concept of a lattice.
Lattice planes

- General Indices: lattice planes are indexed by the inverse of where they cut each axis:
  - Intercept of 0.5 → index=2
  - Intercept of $\infty$ (|| to axis) → index=0

- Related concept: Miller indices
  - used for crystal faces
  - Contain no common factors

- Notation: $[ ]$ defines a direction
  - $[100]$ is along $a$ axis
The Unit Cell

- The **unit cell** describes at least one repeating unit that can be used to construct the structure.
- There are 7 types of unit cells corresponding to the 7 **crystal systems**.

Centering is used to increase symmetry

- The green (primitive) unit cell does not demonstrate two-fold symmetry that can be seen in the red (centered) cell.
Centering causes lattice points to be placed inside units cells (body center, face centers) giving rise the 14 Bravais lattices (1848)

Have non-perpendicular axes: (non-orthogonal coordinate systems)

(Figure from http://www.chemsoc.org/exemplarchem/entries/2003/bristol_cook/latticetypes.htm)
Symmetry

The construction of a crystal from the unit cell requires repeated translation of the “building block” in all three directions: **lattice symmetry**

Additional symmetry is almost always present between the atoms in a unit cell. This means the unit cell (and thus the entire structure) can be built from just a section of the unit cell—The minimal section representative of the entire structure is called the **asymmetric unit**

- Types of symmetry elements in crystals
  - Lattice translations (includes lattice centering)
  - Mirror planes
  - Proper/improper Rotation axes (includes center of symmetry)
  - Screw Axes
  - Glide Planes

(Images from http://members.tripod.com/~EppE/302rev2.htm)
Space Groups

- Not all combinations of symmetry and lattice types are compatible
  - Example: mirror plane perpendicular to a non-orthogonal pair of axes – is not possible

△ There are only 230 unique ways to combine symmetry elements for a 3D lattice: **230 space groups**

△ Space groups are tabulated in *The International Tables of Crystallography, Volume A*! I recommend *Space Groups for Solid State Scientists* by G. Burns and A. M. Glazer as a good place to learn about space groups and s.g. properties
Symmetry, Space groups

- Translational symmetry (centering, screw axes, glide planes) causes classes of reflections to have zero intensity: systematic extinctions
  - Non-translational symmetry (mirror planes, rotation axes, centers of symmetry) do not do this.

- Space groups describe the 230 (219 if chirality differences are ignored) ways that symmetry elements can be combined.
  - Symmetry provides the scaffolding that structures are built upon

Fractional coordinates

- Atom locations are measured in fractions of the unit cell edges
  - Note atom is at $x=0.45, y=0.25$

\[ \triangle \] This notation allows for simple description of symmetry operations:
- $(x,y,z) \rightarrow (1+x, y, z)$ [translation on $x$]
- $(x,y,z) \rightarrow (1/2+x, 1/2+y, 1/2+z)$ [centering]
- $(x,y,z) \rightarrow (-x, -y, -z)$ [center of symmetry @ origin]

In crystallographic notation $x=0.45(3)$ means that there is a standard uncertainty of 0.03 on the value for $x$ of 0.45

Equivalently, there is a 95% chance that $x$ is between 0.39 and 0.51 ($\pm 2\sigma$)
Reciprocal Lattice

- To simplify math when working with non-orthogonal coordinate systems, we use a construct called the **reciprocal lattice** (indicated by star) where each reciprocal axis is perpendicular to two “real space” axes:
  - $a^* \cdot a = 1; \ a^* \cdot b = 0; \ a^* \cdot c = 0$
  - $b^* \cdot a = 0; \ b^* \cdot b = 1; \ b^* \cdot c = 0$
  - $c^* \cdot a = 0; \ c^* \cdot b = 0; \ c^* \cdot c = 1$

- This means that if we have two vectors:
  
  \[ \mathbf{r} = xa + yb + zc \quad \text{and} \quad \mathbf{d}^* = h a^* + k b^* + \ell c^* \]

  Then no cross-terms are needed for the dot product:
  
  \[ \mathbf{r} \cdot \mathbf{d}^* = hx + ky + \ell z \]

*Use of the reciprocal lattice makes computation of the Fourier transform of the atomic positions straightforward.*

*Historical note: the value of the reciprocal lattice for working with non-orthogonal coordinate systems was first recognized by J. Willard Gibbs (1881)*
Where to go for more…

There are many texts available. My favorites:

[Modern & very comprehensive, quite reasonable price considering quality, size & scope.]

[Focused on small-molecule single crystal techniques, dated, but very easy to read; very good explanations of fundamentals. 1st book for many in field.]

**APS Web lectures on powder diffraction crystallography:**  
(http://www.aps.anl.gov/Xray_Science_Division/Powder_Diffraction_Crystallography)  
Intended to introduce Rietveld refinement techniques with GSAS & EXPGUI