

3.2. THE PHYSICS OF DIFFRACTION FROM POWDERS

$$\frac{4 \sin^2 \theta}{\lambda^2} = d^{-2} = Ah^2 + Bk^2 + Cl^2 + Dkl + Ehl + Fhk, \quad (3.2.1)$$

where $A = a^{*2}$, $B = b^{*2}$, $C = c^{*2}$, $D = 2b^*c^* \cos \alpha^*$, $E = 2c^*a^* \cos \beta^*$ and $F = 2a^*b^* \cos \gamma^*$. Crystal symmetries higher than triclinic lead to significant simplifications in the above, e.g. in the orthorhombic system, $a^* = 1/a$, $A = 1/a^2$ etc., $\alpha^* = \beta^* = \gamma^* = 90^\circ$, and $D = E = F = 0$. See Chapter 1.1, *Reciprocal space in crystallography*, in Volume B of *International Tables for Crystallography* for more details.

3.2.2.2. Diffraction peak intensities

3.2.2.2.1. X-rays

Consider a polycrystalline sample of randomly oriented grains of a single crystalline phase, containing a total number of atoms N , illuminated by an X-ray beam of wavelength λ . Each reciprocal-lattice vector with magnitude $|\mathbf{G}| < 2/\lambda$ will produce a cone of diffracted radiation with opening half-angle $2\theta = 2 \sin^{-1}(\lambda|\mathbf{G}|/2)$. Regarding the diffraction lines as perfectly sharp delta-functions, the differential cross section for one powder Bragg peak is (Marshall & Lovesey, 1971)

$$\frac{d\sigma}{d\Omega} = \frac{N\lambda^3 r_e^2}{16\pi V \sin \theta \sin 2\theta} |A_{hkl}|^2 \delta\left(2\theta - 2 \sin^{-1} \frac{\lambda}{2d}\right). \quad (3.2.2)$$

In this equation, $r_e = 2.82 \times 10^{-15}$ m is the classical electron radius and V is the unit-cell volume; the polarization factor P , multiplicity m_{hkl} and structure factor A_{hkl} are discussed below. Finite crystallite size, strain and other factors discussed below will broaden the peak, but the integrated intensity of each peak will be given by the factors before the delta-function.

Before continuing, note that this result is correct within the approximation that the entire sample is uniformly bathed in radiation (i.e., there is no absorption) and the kinematic approximation that the scattering power is very weak, so that further interaction of the scattered radiation with the sample may be neglected. The latter is equivalent to the Born approximation of elementary quantum mechanics, and its failure is referred to as extinction; it is discussed below in Section 3.2.3.6. It should also be noted here that equation (3.2.2) refers only to diffraction in Bragg peaks, and not to thermal diffuse scattering or scattered radiation due to static distortions from an ideal crystal lattice (Huang scattering; Krivoglaз, 1969).

Returning to the terms in equation (3.2.2), for X-rays the polarization factor P accounts for the polarization dependence of the Thomson scattering cross section from a free electron. For a completely polarized incident beam: $P = 1$ for S polarization (polarization perpendicular to the scattering plane); $P = \cos^2 2\theta$ for P polarization (in the scattering plane). For unpolarized X-rays (no monochromator), $P = (1 + \cos^2 2\theta)/2$. For horizontally polarized X-rays from a synchrotron-radiation source, diffracted vertically, the polarization factor is very close to unity, and so P can generally be taken as 1. If there is a crystal monochromator (deflection angle $2\theta_m$) coplanar with the diffractometer and an unpolarized source, $P = (1 + \cos^2 2\theta_m \cos^2 2\theta)/(1 + \cos^2 2\theta_m)$. If there is a polarization-sensitive analyser crystal following the sample, its relative transmission of X-rays polarized within or perpendicular to its plane of scattering must also be taken into account.

The multiplicity factor m_{hkl} accounts for the number of symmetry-equivalent reflections in the crystal structure, i.e., the number of distinct reflections that have the same structure factor. Note that the equivalence of Bragg reflections depends on the

Laue group of the sample. For example, in the cubic space group $Fm\bar{3}m$, reflections (hkl) and (khl) are equivalent, whereas they are not in space group $Fm\bar{3}$. If the atomic scattering factors are purely real or the structure is centrosymmetric, the intensities of Friedel pairs hkl and $\bar{h}\bar{k}\bar{l}$ are equal. However, in an acentric structure with significant imaginary component of any of the atomic scattering factors (see below), Friedel pairs cannot be treated as symmetry equivalent.

The (dimensionless) X-ray structure factor is given by

$$\begin{aligned} A_{hkl} &= \sum_n f_n \exp(2\pi i \mathbf{G}_{hkl} \cdot \mathbf{r}_n) \exp(-W_n) \\ &= \sum_n f_n \exp[2\pi i(hx_n + ky_n + lz_n)] \exp(-W_n), \end{aligned} \quad (3.2.3)$$

where the sum runs over all atoms in the unit cell. In the first form of this equation, \mathbf{r}_n is the vector from the origin of the unit cell to the n th atom; in the second form, x_n , y_n and z_n are fractional coordinates of the n th atom in the unit cell such that $\mathbf{r}_n = x_n \mathbf{a} + y_n \mathbf{b} + z_n \mathbf{c}$.

The term $\exp(-W_n)$ is called the Debye–Waller factor, and it accounts for random dynamical motion of the n th atom away from its equilibrium position. This may be due to thermal fluctuations or (at low temperature) the quantum zero-point motion. If the atomic motion is isotropic, this term takes the form

$$W_n = B \sin^2 \theta / \lambda^2 = 8\pi^2 U_n \sin^2 \theta / \lambda^2, \quad (3.2.4)$$

where $U_n = \langle u_n^2 \rangle$ is the mean-square fluctuation of the atom's distance from its equilibrium position (in three dimensions). Observed values of B are generally in the range of 0.5 to 1 Å² for inorganic crystals, and as large as 5 Å² for organic crystals.

A more general treatment of the Debye–Waller factor considers the probability density of the centre of the atom to be a three-dimensional ellipsoidal Gaussian function. Then for a given diffraction peak, it takes the form

$$\begin{aligned} W &= 2\pi^2(U_{11}a^{*2}h^2 + U_{22}b^{*2}k^2 + U_{33}c^{*2}l^2 \\ &\quad + 2U_{12}a^*b^*hk + 2U_{23}b^*c^*kl + 2U_{13}a^*c^*hl). \end{aligned} \quad (3.2.5)$$

The parameters U_{ij} that define the displacement ellipsoid are constrained to match the point symmetry of the site. For example, an atom that lies on a site of mmm symmetry will have the cross terms $U_{12} = U_{23} = U_{13} = 0$.

In equation (3.2.3), f_n is the atomic scattering factor, which arises because X-rays interact with the electrons in the sample, and the strength of that interaction depends both on the magnitude of the scattering vector and the X-ray frequency (wavelength). The electrons are distributed around the nucleus, and so their scattering power decreases with increasing magnitude of the scattering vector. There is also a dependence of the phase and amplitude of the scattering factor on the X-ray energy, which is especially significant near resonance with atomic transition energies. These factors are gathered into the atomic scattering factor f_n , which is commonly written as

$$f_n = f_n^0(\sin \theta / \lambda) + f'(\lambda) + if''(\lambda). \quad (3.2.6)$$

f^0 is the Fourier transform of the electron number density, so that $f^0(0)$ is equal to the number of electrons in the atom or ion. These factors are tabulated in *International Tables for Crystallography*, Volume C, Chapter 6.1 and <https://physics.nist.gov/PhysRefData/FFast/html/form.html> and http://henke.lbl.gov/optical_constants/asf.html. Values of atomic scattering factors are normally incorporated into data-analysis programs that require them, so the user does not often have to worry about them.

3. METHODOLOGY

Equation (3.2.3) above is premised on the assumption that the distribution of electrons in the sample is confined to spherically symmetric atoms, *i.e.*, ignoring charge density in bonds and the deformation of core electron density. For the purposes of phase identification and structure refinement, this is generally a good approximation. X-ray determination of non-spherical charge densities goes beyond the scope of this introduction; for further information, see *e.g.* Coppens (1997) and Bindzus *et al.* (2014).

In energy-dispersive X-ray measurements, the detector sits at a fixed 2θ and collects diffracted radiation from a continuum source. The independent variable of the measurement can be taken as the X-ray energy, $E = hc/\lambda$. With the appropriate change of variables, equation (3.2.2) can be written as

$$\frac{d\sigma}{d\Omega} = \frac{Nr_e^2}{16\pi V} \frac{h^3 c^3 P m_{hkl}}{E^2 \sin^3 \theta} |A_{hkl}|^2 \delta\left(E - \frac{hc}{2d \sin \theta}\right). \quad (3.2.7)$$

Recall that $hc = 12.398 \text{ KeV \AA}$ in convenient units. Again, instrument resolution and sample effects will broaden the peak, but its integrated area is given by the terms preceding the delta-function. Of course, the diffracted intensity must be normalized to the incident spectrum as a function of energy; this may be a rather complicated undertaking as it involves energy-dependent corrections for such factors as detector sensitivity, absorption in the sample *etc.*

3.2.2.2.2. Neutrons and nuclear scattering

Neutrons interact with the sample in two ways: the strong interaction with the atomic nuclei, and the magnetic interaction between the neutron's dipole moment and magnetization density in the sample. We consider here only unpolarized neutrons; the use of polarized neutrons permits separation between nuclear and magnetic scattering as well as direct observation of the interference between the two; details are beyond the scope of this chapter. Nuclear scattering is very similar to the X-ray case discussed above, except that the atomic scattering amplitude $r_e f_n$ is replaced by the nuclear coherent scattering length b_n (given in *International Tables for Crystallography*, Volume C, Table 4.4.4.1), which is generally independent of neutron energy.

Unlike X-rays, the strength of the neutron–nucleus interaction is not a smooth function of atomic number. This creates opportunities to use neutrons to distinguish atoms with nearly identical X-ray scattering amplitudes, but it also makes certain elements very difficult to study with neutrons. The interaction between neutrons and the nuclei in the sample depends on the isotope and possibly the spin angular momentum of the neutron–nucleus system. This means that incoherent scattering can be significantly larger than the (coherent) diffracted signal for certain atoms, notably hydrogen (^1H); see Chapter 2.3 of this volume for further details. For wavelengths of interest in crystallography, the nucleus is essentially a point, and so there is no atomic form factor. This generally leads to greater intensity relative to X-rays at increasing scattering vector (decreasing d -spacing).

Neutron diffractometers operate in one of two ways: angle dispersive or energy dispersive. The configuration for angle-dispersive diffraction measurements is conceptually similar to that used for X-rays; a monochromatic beam of neutrons impinges on the sample and a detector measures the distribution of neutrons *versus* scattering angle. For Bragg neutron diffraction from nuclei,

$$\frac{d\sigma}{d\Omega} = \frac{N\lambda^3}{16\pi V \sin \theta \sin 2\theta} \frac{m_{hkl}}{|A_{hkl}|^2} \delta\left(2\theta - 2\sin^{-1} \frac{\lambda}{2d}\right), \quad (3.2.8)$$

where the neutron nuclear structure factor is defined as

$$A_{hkl}^{(n)} = \sum_n \bar{b}_n \exp(2\pi i \mathbf{G} \cdot \mathbf{r}_n) \exp(-W). \quad (3.2.9)$$

Time-of-flight neutron diffractometers, generally based at pulsed spallation sources, operate by measuring the time from the creation of the pulse of neutrons at the target until they appear in a given detector. If the total path length from source to detector is L and the detector is situated at an angle 2θ , a neutron with time of flight t had speed L/t and wavelength $\lambda = ht/m_n L$. Here h is Planck's constant and m_n is the mass of the neutron. This provides a measurement of the d -spacing within the sample, $d = ht/(2m_n L \sin \theta)$. Another change of variables from equation (3.2.2) yields

$$\frac{d\sigma}{d\Omega} = \frac{N}{64\pi V} \left(\frac{h}{m_n L}\right)^3 \frac{t^4}{\sin^2 \theta} m_{hkl} |A_{hkl}|^2 \delta(t - 2d \sin \theta L m_n / h). \quad (3.2.10)$$

In practice, a large number of detectors surround the sample and counts from the same d -spacing (appropriately normalized for incident-beam intensity and detector solid angle) are binned together. In convenient units, $m_n/h = 253 \mu\text{s m}^{-1} \text{ \AA}^{-1}$.

3.2.2.2.3. Neutrons and magnetic scattering

Magnetic neutron scattering is also described through a structure factor which is, however, a vector. The magnetic moment of the neutron interacts with the magnetization density of unpaired electrons in the sample, which may possess spin and/or orbital angular momentum. The magnetic interaction is only sensitive to the component of magnetization perpendicular to the scattering vector. When discussing magnetic scattering, it is more common to use the scattering vector $\mathbf{Q} = 2\pi\mathbf{G}$. The magnetic structure factor is defined as

$$\mathbf{A}_{\mathbf{Q}}^{\text{mag}} = (\gamma r_e / 2) \sum_n f_n(\mathbf{Q}) (\hat{\mathbf{Q}} \times \mathbf{m}_n \times \hat{\mathbf{Q}}) \exp(i\mathbf{Q} \cdot \mathbf{r}_n) \exp(-W). \quad (3.2.11)$$

Here $\gamma = 1.9132$ is the neutron gyromagnetic factor, $f_n(\mathbf{Q})$ is the atomic magnetic form factor, \mathbf{m}_n is the magnetization of the n th site in units of the Bohr magneton and $\hat{\mathbf{Q}}$ is the unit vector in the direction of \mathbf{Q} . The double cross product isolates the component of magnetization perpendicular to \mathbf{Q} .

Note that many magnetically ordered materials have a magnetic cell which is larger than the chemical cell. Indeed, many magnetic phases are incommensurately modulated, *i.e.*, the magnetic structure is not periodic with any combination of the chemical unit cell translation vectors. Such matters are beyond the scope of this introduction, and are handled in Chapter 7.13 of this volume.

For unpolarized neutron measurements (*i.e.*, an average over all polarization states of the incoming and diffracted beam), the intensities of the nuclear and magnetic diffraction peaks may be computed separately and then added to determine the overall diffraction pattern. In cases where the chemical and magnetic cells are identical (*e.g.* simple ferromagnets) the nuclear and magnetic diffraction patterns overlap, and so one observes only intensity differences upon magnetic ordering. In the case of antiferromagnets, new magnetic diffraction peaks appear at positions not allowed for the chemical unit cell.

Note also that the magnetic form factor depends on the spin density in the magnetic orbitals, which are typically of greater spatial extent than either the total charge density or the nuclear