12.1-01 NUMERICAL EVALUATION OF INTENSITY ABERRATIONS. By J. H. Konnert and P. D'Antonio, Laboratory for the Structure of Matter, Naval Research Laboratory, Washington, D. C. 20375, U.S.A.

It is important in the collection and interpretation of diffraction data to evaluate the aberrations affecting the position, shape and intensity of details in the diffraction pattern. Among contributing factors are: source size and shape; incident beam collimation; size, shape, position and absorptivity of sample; diffracted beam collimation; monochromater (if any) efficiency, and angle setting errors. For many experimental arrangements, it is not possible to express analytically the cumulative effects of these factors. However, modern computers permit the numer-

ical evaluation of aberrations for arbitrary source, collimation and detector configurations. We have found useful a computer program that calculates for a set of experimental conditions the "profiles" that indicate the shape, position and intensity of curves that, for perfect resolution, would be delta functions. The first figure illustrates the method by which these profiles are calculated. The profile illustrated derives from a single diffraction cone of $2\theta = 1^{\circ}$ determined by a single point on the source, a single point in the sample, and a detector slit. As an



insert in this figure, the geometry in the detector slit plane is also illustrated. The approximate circle represents the intersection of the diffraction cone with this plane. The detector slit may be thought of as sweeping from the center of the cone to the right as the detector ranges from $2\theta = 0^{\circ}$ to 1° . In this example, a monochroranges from $2\theta = 0^{\circ}$ to 1° . In this example, a monochromater is not present, and the observed intensity is proportional to the fraction of the cone passing through the detector slit. At 0.56° , the intensity rises abruptly as the 2nd portion of the cone also passes through the slit. At $\sim 1^\circ$, a long cone length passes through the slit resulting in the sharp peak. Complete profiles may be computed by considering the many diffraction cones arising from finite source and sample size. While the simple geometry represented here may be express-ed analytically, incident beams and diffraction cones may be evaluated numerically for the passage through any collimation system. The second figure illustrates representative profiles for slit (Kratky Camera) and approximate pinhole geometries. These low angle profiles have been chosen for illustrative purposes because of the large associated aberrations. Higher angle data will

generally be affected by smaller aberrations. Recognition of such small aberrations, however, may be quite important when carrying out analyses such as profile refinements. Aberrations associated with several experimental geometries will be illustrated along with details of the numerical evaluations of the profiles.



12.1-02 AN AUTOMATED POWDER DIFFRACTION DATA REDUCTION PROGRAM FOR LARGE THROUGHPUT. By <u>G.</u> <u>R. Fischer</u> and W. T. Kane, Corning Glass Works, R&D Division, Corning, New York - 14830

A fully automated X-ray powder diffraction system at Corning Glass Works was specifically designed for large throughput of 35 or more samples per day. The required operator time is drastically reduced because functions such as sample-log keeping, running of the diffractometer and sample changing, and the numbering and labeling of the patterns and individual peaks are performed by computer. The data reduction part of the system copes automatically and without operator interaction with a large variety of materials ranging from fully amorphous to highly crystalline specimens.

Several novel approaches in the data reduction process were required to achieve the versatility needed. These include the filtering of the (fast) Fourier transform of the data set with an auto-adjusting digital filter, the approximation of the general background through the fitting of spline functions, and the automatic calculation of a variable threshold for peak acceptance based on a pseudo-standard deviation.

The program is written in a high level language (FORTRAN), easily transportable, and executes rapidly even on moderate sized minicomputers. The output consists of a table of 20 (corr) d (corr), and I/I_1 and of a plot of the refined X-ray pattern with all peaks labeled with their corresponding d (corr) and observed relative intensities.

12.1-03 EXPERIMENTAL & COMPUTATIONAL TECHNIQUES FOR RECORDING & ANALYSING PHOTOGRAPHIC POWDER DATA. By <u>M.J.Mendelssohn</u> & H.J.Milledge (Crystallography Unit, Department of Geology) & D.Walley (Computer Science Department), University College London, Gower Street, London WC1E 6BT, England.

During investigations designed to optimise the parameters involved whenfluorescent screens are used to register diffraction data, experiments are being made using radiation selected from the continuum (a situation which arises in both neutron and in synchrotron diffraction experiments). The tests involving powder samples are being carried out using a Stoe powder camera which permits both transmission and reflexion patterns to be recorded with monochromatised radiation.

Since the efficiency of fluorescent screens increases very rapidly as the wavelength decreases, patterns produced with radiation of $\sim 0.5 - 0.7A$ (near the peak of the continuum) can appear to be considerably stronger than those obtainable with the K \propto radiation from, for example, a Cu tube. Thus, if adequate resolution can be achieved, the continuum radiation may actually be preferable for some purposes, especially those involving real-time phenomena such as phase transitions, encapsulated specimens where hard radiation can reduce effects due to absorption, or situations where anomalous dispersion effects may be maximised by the choice of a particular wavelength, as in the Laue case (Grenville-Wells & Lonsdale, Nature (1954)<u>173</u> 1145).

In order to evaluate these non-standard data sets they are being compared with conventional powder patterns obtained from the same specimens and measured with the same computer-controlled scanning microdensitometer.