PREPARATION OF NUCLEAR-TRACK PLATES AND STRIPPING FILMS FOR THE STUDY OF RADIOACTIVE MINERALS*

L. R. STIEFF AND T. W. STERN,

U. S. Geological Survey, Washington, D. C.

Abstract

Some modifications of current techniques for preparing nuclear plates for mineralogical studies are described. A method of mounting nuclear-track stripping film on thin sections is given. Illustrations are presented of the application of these techniques to the study of the radioactivity of powdered rocks, minerals, and thin sections.

INTRODUCTION

Since Becquerel's discovery of radioactivity in 1896 by the exposure of an unactivated uranyl sulfate phosphor to a photographic emulsion, the photographic method has become a valuable tool in the study of radioactive minerals. Autoradiographic plates, nuclear-track plates, and autoradiographic stripping film may be used to recognize radioactive minerals and to study the distribution of radioactivity in rocks. In addition photographic emulsions have been used in many specialized mineralogic studies, not discussed in this paper, such as the semiquantitative determination of both uranium and thorium in the same rocks or minerals (Coppens, 1950, p. 21; Picciotto, 1949, p. 80; Yagoda, 1949, p. 160), the departure of uranium minerals from radioactive equilibrium (Yagoda, 1949, pp. 101, 182), the artificial autoradiography of minerals (Goodman and Thompson, 1943, p. 456), and the natural fission rates and neutron fluxes in uranium ores (Yagoda and Kaplan, 1949, p. 702).

The autoradiographic method of studying the distribution of radioactivity in rocks is the oldest and simplest. A polished rock or mineral section is placed in contact with the emulsion of a photographic plate sensitive to either light, to x-rays, or to alpha particles. After the plate has been properly exposed and developed, the radioactive minerals are recognized by comparing the pattern on the photographic plate with the polished specimen. Bowie (1951, p. 65) and Poole and Bremner (1949, p. 130) using mechanical means of registering the autoradiographic plate and specimen have been able to locate radioactive grains 20 to 25 microns in diameter. If more precise registration is required, however, mechanical methods of alignment are not entirely satisfactory. The microscopic study of the superimposed alpha plate is also limited to microscope objectives which have greater working distances than the

* Publication authorized by the Director, U. S. Geological Survey.

thickness of the glass autoradiographic plate. Because of these shortcomings and because this simple method has been fully described by Yagoda (1946*a*, p. 87) and others, no further elaboration of this technique is given herein.

The dispersed-grain method of Mügge (1909, p. 114), of Tyler and Marais (1941, p. 146), and of Yagoda (1949, p. 176) has solved the problem of registering the developed alpha or beta tracks with the radioactive grain that emitted the energized particle, by entrapping powdered rock or mineral fragments in a water-softened emulsion. This method uses special photographic plates covered with thick, fine-grained emulsions (nuclear emulsions) capable of recording the trajectories of alpha and beta particles emitted by both natural and artificial radioactive elements. Although this technique provides a means of recognizing radioactive grains, obviously it can give little information on the spatial distribution of the radioactive minerals in the rock. In spite of this drawback, this technique has many important uses, and the modifications of it developed in the Trace Elements Section, Washington Laboratory of the Geological Survey are described below.

The stripping-film method has been used in tracer studies of histological thin sections by Pelc (1947, p. 749), Boyd and Williams (1948, p. 225), MacDonald, Cobb, Solomon, and Steinberg (1949, p. 117), Berriman, Herz, and Stevens (1950, p. 472), Doniach and Pelc (1950, p. 184), and Heller (1950, p. 8), but, to our knowledge only Bowie (Personal communication, 1951) has used stripping film for geologic studies. The use of stripping film permits positive recognition of radioactive grains, and the permanent registration of the emulsion and the thin section also provides a means of studying the spatial distribution of the radioactivity in the rock. We have successfully used the method described below in mounting stripping films on more than 100 thin sections of rock.

DISPERSED-GRAIN METHOD

Type of plate.—The alpha-sensitive nuclear-track plates used in the dispersed-grain method may be obtained from the Eastman Kodak Co., of Rochester, N. Y., and from Ilford Ltd., and Kodak Ltd., in England. Eastman and Ilford emulsions used for this work are designated as NTA and C-2, respectively. These plates are available in various sizes from 1 by 3 inches to 8 by 10 inches. Emulsions are available in thicknesses of 25, 50, 100, and 200 microns. An emulsion thickness of 50 microns is recommended for mineralogical work because the maximum range of the most energetic alpha particles in the emulsion does not exceed 50 microns. Sixty days should be allowed for the delivery of these plates.

For this method Kodak NTA plates $1\frac{1}{4}$ by 10 inches with an emulsion thickness of 50 microns were used. These plates are sensitive to light and should be handled under a $7\frac{1}{2}$ -watt safelight with a Wratten series OA filter. The plates, emulsion side down, are cut into $1\frac{3}{4}$ -inch lengths by means of a small plate-cutting jig shown in Fig. 1. A Goodell-Pratt wheel-type glass cutter is used to score the glass side of the plate. After the plate is scored, it is advanced and broken over the end of the jig.



FIG. 1. Alpha-plate cutting jig.

An identifying number is then inscribed on the glass side of the plate with a diamond-pointed pencil.

Before studying rocks containing trace amounts of uranium and thorium, it may be desirable first to remove from the plate the tracks that have accumulated since the plate was manufactured. These tracks are due to traces of uranium and thorium in the emulsion. Yagoda(1946*a*, p. 94; Wiener and Yagoda, 1950, p. 39) has found that the background tracks may be eradicated by placing the plates in a light-tight container over a 3-per cent solution of hydrogen peroxide for 4 hours.

Sample preparation.—Usually specimens to be studied by this method should be ground and sized. Material of sieve sizes from minus 80 to minus 200 mesh has been used with success. If necessary, the sample may be further separated by magnetic or heavy-liquid methods and the resulting separates placed on a nuclear-track plate (Fig. 2). Also chemi-

cal separation of the sized material before dispersal is possible. After the sample is prepared, the sized fractions may be stored in a closed container for approximately a month to allow radon and its daughter products in the sample to return to radioactive equilibrium. Prior to preparing the plates, approximately 20 mg. of this sized material is dispersed in 10 ml. of distilled water in a capped 50-ml. bottle.



FIG. 2. Photomicrograph of crystalline carnotite from the May Day mine, Mesa County, Colo., dispersed on a nuclear-track plate. The tracks in the emulsion diverging from the carnotite grains are the paths taken by the alpha particles emitted by uranium or its radioactive daughter products. The time of alpha-plate exposure was 5 days.

Plate preparation.—A 6-cm. petri dish is filled with 10 ml. of distilled water at room temperature. The $1\frac{3}{4}$ -inch alpha plate is placed, emulsion side up, in the petri dish and allowed to soak for 5 minutes. The emulsion has then softened sufficiently to trap the dispersed material which now is added near the edge of the petri dish. Care must be taken not to pour the dispersed material directly over the plate, as this loads it unevenly. The plate will now be covered with approximately 8 mm. of water. The time required for the sample to settle out depends primarily on the distribution of particle sizes in the sample and can be calculated from Stokes' law. A settling time of 5 minutes is usually sufficient for all but the finest-sized samples. The plate, now being loaded with approximately 0.007 mg. of sample per square millimeter, is carefully removed from the petri dish with tweezers and placed to dry, emulsion side up, on a flat surface. When the plate has dried, it should be stored in a light-tight box for exposure. If prolonged exposure (1,000 hours or more) is neces-

sary, surface reactions and fermentation of the gelatin may be minimized by placing the box containing the plate in a refrigerator (Picciotto, 1949, p. 77).

Exposure.—The length of exposure is related to the following: the uranium or thorium content of the sample, the degree to which the uranium or thorium is in radioactive equilibrium with its daughter products, the size and shape of the grains, and the manner in which the radioactivity is distributed in the sample. Generally, a minimum of two diverging alpha tracks is necessary to establish the radioactivity of a grain. Small pitchblende grains can be recognized easily after a 72-hour exposure, but the extremely fine grained radioactive segregations in phosphate rocks require an exposure of at least 1,000 hours. If the length of exposure required for satisfactory results with a mineral grain is unknown, several plates should be prepared at the same time and exposed for different periods of time. Yagoda (1949, p. 177) has calculated that "a fragment measuring 10 microns in average diameter, containing about 50 per cent uranium, will record about 20 tracks per day of exposure. The isolated grain contains less than 10^{-8} g. of uranium"

Development.—After exposure the plate is developed from 2 to 4 minutes at 68° F. in full strength Eastman Kodak D-19 developer (or Ilford, ID-19). This type of development shows the alpha particle energy-loss rate by changes in the grain spacing of the track. However, if only track counts are to be made, the plate should be developed for 2 minutes at 68° F. in Eastman Kodak D-8 developer diluted two to one (Webb, 1948, p. 519). Alternate methods of development are given by Yagoda (1949, p. 60). The plate is then rinsed for 10 minutes with two changes of water. The plate is fixed in Eastman Kodak F-5 fixer or 30 per cent hypo for twice the time required to clear the emulsion and rinsed for 15 minutes with three changes of water. Finally, the plate is allowed to drain and dry thoroughly in a dust-free atmosphere.

Plate covering.—The dispersed-grain methods of Tyler and Marais and others do not include covering the alpha plates. We have found, however, that the addition of a cover glass serves two purposes. First, the alpha plate becomes a permanent mount and may be used repeatedly without danger of dislodging the grains. Second, as most microscope objectives are designed to be used with covered specimens and as alpha plates generally are studied under high-power magnification, the use of a cover glass results in a marked improvement in the resolution of the tracks and mineral grains.

A clean glass cover slip, 22 mm. square, is mounted on the alpha plate with two drops of Canada balsam in xylol. The plate is now kept in an oven in a horizontal position for 48 hours at a temperature of approxi-

188

mately 140° F. After removal from the oven, the plate is allowed to cool before it is examined or stored.

Microscopic examination.—The alpha tracks emitted by the radioactive grains are studied by focusing the microscope down into the emulsion (Fig. 2). Although some alpha tracks may be seen at magnifications of approximately 100 diameters, quantitative track counts cannot be made successfully at magnifications of much less than 500 diameters. Detailed studies of track length and grain spacing should be made at magnifications in excess of 1,000 diameters.

For rapid surveys of the alpha plates a binocular metallographic microscope is used. These surveys are made with transmitted light, a $40 \times$ objective, and $15 \times$ planar oculars. A $95 \times$ oil-immersion objective (N.A.=1.25) is used with the same oculars for detailed track and mineral studies. A larger field can be obtained by using a Bausch and Lomb $40 \times$ fluorite oil-immersion objective (N.A.=1.00). For identification of the radioactive minerals a petrographic microscope is employed.

If track-count studies are to be made on previously identified mineral grains, eyestrain can be reduced by the use of a green monochromatic filter or by dark-field illumination.

Advantages and limitations.—The dispersed-grain method may be used to recognize the radioactivity of minerals in a powdered rock sample and may aid in the identification of the minerals. If alpha plates are made under controlled conditions, departures from radioactive equilibrium and the approximate uranium and thorium content may be estimated. A quantitative determination of the uranium content of radioactive minerals by an analysis of the tracks is laborious and time consuming, but a qualitative estimate of the radioactivity of minerals may be made much more easily. Radiocolloids, the localized concentrations of shortlived daughter products (Yagoda, 1946b, pp. 462–470), may be studied easily. Alpha plates also may aid in the selection of samples for leaduranium-ratio age determinations.

This method is limited to minerals that are not water soluble and that are not attacked by photographic solutions. Also, this method can give little information on the spatial distribution of the radioactive minerals in the rock.

STRIPPING-FILM METHOD

Type of film.—Kodak autoradiographic stripping film consists of a 150-micron supporting film to which has been bonded a 10-micron cellulose ester film covered with a 10- or 25-micron nuclear emulsion sensitive to alpha and beta particles (NTB). (See Fig. 3A.) These NTB stripping films may be obtained from the Eastman Kodak Co., of Rochester, N. Y. Stripping film is at present available only in boxes containing ten 4-by-5 inch sheets. Approximately 60 days should be allowed for delivery of this film.

Film preparation.—NTB emulsions are sensitive to light and should be handled before development under a 25-watt safelight equipped with a Wratten series 2 filter. A sheet of stripping film, measuring 4 by 5 inches, is cut with a print trimmer into eight pieces, each 1 by $2\frac{1}{2}$ inches.

Two score marks, parallel to the length and one-eighth of an inch from each edge, are made through the emulsion and cellulose ester film with a



FIG. 3. Nuclear-track stripping film across section and preparation. (A) Cross section. (B) Film, emulsion side up. (C) Film, supporting base up. (D) Unsupported emulsion.

razor blade and straightedge. (See Fig. 3B.) As the film is pressure sensitive, care must be taken to avoid touching the part of the emulsion between the score marks. The razor should not cut through the 150micron supporting base. We have found that the tendency of the film to tear during the stripping process is minimized if the film has been scored as described above.

Approximately half an inch of one end of the film then is placed, emulsion side down, on a small wooden block and a score 1 inch long and approximately a quarter of an inch from the end of the piece of film is made with a razor blade and straightedge (Fig. 3C). This score is deeper than the $2\frac{1}{2}$ -inch score marks and should go about halfway through the supporting base. A similar score is made on the other end of the film.

190

The film, removed from the wooden block, is held by the edges, emulsion side down, and the supporting base is broken carefully along the 1-inch score by bending the tab toward the emulsion. This operation is repeated for the other end of the film. With the film held in the left hand and the tab in the right hand, gently start to strip the emulsion and the cellulose ester film from the supporting base. The emulsion and cellulose ester film should be freed from the supporting base all along the 1-inch score along one end of the film before this operation is repeated for the other end. Now, holding the film with the left hand, emulsion side toward the operator, slowly pull the tab, freeing the emulsion and cellulose ester film from the supporting base. If the film begins to tear, the tear usually will not cross the lengthwise score marks. If the film is stripped without difficulty, the $\frac{1}{8}$ -inch ribbons can be removed with tweezers. (See Fig. 3D.) It is essential that the film be stripped slowly to prevent fogging by static electricity. It has been found that the film can be stripped more easily on days of high relative humidity. When the relative humidity is low, the film should be placed in a box with a piece of moistened blotting paper for about 15 minutes before the stripping operation is started.

Stripping-film jig.—The jig used to hold the stripping film during mounting is shown in Fig. 4A. This jig may be made of metal, wood, or glass slides; its dimensions depend on the size of the specimen to be covered. A rubber band, not shown in Fig. 4, holds the two tab clamps firmly closed. Tweezers are used to place the tabs in the tab clamps of the jig. The loop of film is mounted in the jig with the cellulose ester side up. The loop is centered in the jig and, if necessary, adjusted by means of the tabs until the reflection of the safelight across the top of the loop is parallel to the ends of film jig slide supports. In this position the stripping film is ready to be mounted on the thin section (Fig. 4B).

Preparation of thin sections.—Thin sections may be prepared in the usual manner but with the cover glass omitted; however, special precautions against contamination must be taken with samples containing small amounts of uranium. The use of disposable "metallographic" emery papers to minimize contamination has been described by Yagoda (1949, p. 39). The rock slice may be mounted on the glass slide with either canada balsam or Lakeside 70 cement.

In order to allow the thin section of the sample to return to its original radioactive-equilibrium condition and to minimize the loss of daughter products by emanation from hydrous minerals, the thin section should be stored for approximately a month in a desiccator.

Covering thin section.—The uncovered thin section is first dipped in a solution of gelatin at approximately 150° F. This gelatin solution is prepared by placing 12 g. of gelatin (Eastman Kodak flake gelatin, item no. *P1099*) in 150 ml. of distilled water, allowing the gelatin to hydrate

for 30 minutes, adding 2.5 ml. of 2 per cent chrome alum and 5 ml. of 0.2 per cent aqueous wetting agent (Yagoda, 1949, p. 95) such as aerosol. The solution then should be gently heated. The gelatin is kept in a tightly stoppered wide-mouth bottle on a hot plate except when the slide is being immersed. When not in use, the gelatin solution is stored in a refrigerator to inhibit bacterial growth. To minimize contamination, very low grade samples should not be dipped in the same gelatin solution that is used for high-grade samples.



FIG. 4. Jig for mounting nuclear-track stripping film. (A) Film jig. (B) Film mounted in jig. (C) Slide resting on slide supports.

After dipping, the slide is allowed to drain for approximately a minute to remove excess gelatin. It is ready to be covered when clear finger-print impressions can be made in the gelatin film on the back of the slide. The slide, rock section down, is centered over the loop of film and gently lowered until the slide rests on the film jig slide supports. (See Fig. 4C.)

The slide is chilled by placing a small ice-filled bottle or chilled metal bar on the slide. This rapidly sets the gelatin and prevents the film from pulling away from the thin section during the next operation. The film is cut with a scalpel along the tab fold lines, freeing the film loop from the tabs. After the ice-filled bottle is removed, the slide is taken from the jig. It is placed, thin section up, on the table, and the excess film is

removed with a razor and straightedge. If for any reason the slide has been covered imperfectly, the stripping film may be removed by soaking the slide in warm water. A solution of Eastman Kodak rapid mounting cement diluted one to three with ethyl acetate is used to paint down and secure the edges of the film to the slide. The slide is then stored in a lighttight box for exposure.

If the operator is not familiar with stripping film, it is suggested that he make several practice mounts in daylight before attempting this method under darkroom conditions.

Exposure.—The same general factors governing exposure discussed in the dispersed-grain method apply to the stripping-film method. It should be mentioned, however, that, because of the relatively small amounts of material represented by a thin section of a radioactive grain, the exposure time may be increased by a factor of three or four over the exposure time used for the dispersed-grain method. Alpha counts of a uniform, randomly chosen area of the thin section have been made to provide a rough estimate of the exposure time. The optimum exposure time of any particular type of material can best be determined, however, by the exposure of several slides of the same material. If for any reason the exposure was incorrect and another thin section cannot be obtained, the stripping film may be removed from the thin section. This is accomplished by first gently rubbing the cemented edges with a cotton swab dipped in ethyl acetate and then placing the slide in warm (110° F.) water for about 10 minutes.

Development.—The development recommended above for nucleartrack plates may also be used for NTB stripping films. Movement of the stripping film due to shrinkage of the emulsion during development and fixation has not been noticed.

Microscopic examination.—In contrast with the dispersed-grain method, the alpha tracks emitted by radioactive grains are studied by focusing up into the emulsion. (See Fig. 5A and B.) The combination of objectives and oculars used on alpha plates may be used with equal success on stripping film. For oil-immersion work a drop of immersion oil is placed directly on the developed emulsion. The oil is removed by wiping the slide gently with a soft absorbent cloth or tissue. To remove the last traces of the oil, the emulsion is cleaned with a drop of xylol on an absorbent tissue. Because the cellulose ester is isotropic, the determination of the optical properties of the mineral grains in the thin section is made without difficulty.

The use of a vertical illuminator with a metallographic microscope has aided the study of alpha tracks that have been emitted directly over opaque grains. The peripheral tracks emitted by opaque grains can be studied easily by transmitted light.

L. R. STIEFF AND T. W. STERN

Advantages and limitations.—The stripping-film method represents an improvement over the mechanical methods of registering autoradiographic plates and thin sections (Poole and Bremner, 1949, p. 130), the method of correlating track patterns with grain shape (Coppens, 1950, p. 21), and the method of simultaneously exposing to light both the alpha plate and the thin section (Picciotto, 1949, p. 79). The use of stripping film assures a positive registration between the developed film and the radioactive grains. Studies have been made that suggest that



FIG. 54. A photomicrograph of carnotite-bearing sandstone from the Rock Creek area, Montrose County, Colo. The white and gray fractured grains are quartz and the interstitial areas contain microcrystalline carnotite. Crossed nicols.

the cellulose ester film protects the thin section from chemical attack by either the developer or the hypo and minimizes the loss of daughter products by emanation. It should be mentioned that great dexterity is not required to cover thin sections with nuclear-track stripping film. After several trials in daylight the average operator can make satisfactory mounts under darkroom conditions.

Some of the inherent limitations of the stripping-film method are a consequence of the methods of preparing the thin sections. Radioactive minerals may be lost easily in grinding; and, if precautions are not taken, the slide may be contaminated by the re-use of grinding powders and rouges. A thorough study of the distribution of radioactivity in a rock can be made only by examining many thin sections. Approximately a

week's exposure is required before any results can be obtained. The stripping-film method is also limited by the activity of the material under investigation. Generally, two alpha tracks are necessary to locate a radioactive grain because the alpha tracks must be projected through the 10-micron cellulose ester film. Also quantitative track length studies are not possible because only part of the total track is recorded in the 10micron emulsion and some alpha particle energy is lost in traversing the 10-micron cellulose ester film.



FIG. 5B. A photomicrograph of the alpha tracks overlying the interstitial carnotite in the field shown in figure 5A. The exposure time for the nuclear-track stripping film was 6 days. Crossed nicols.

Recently Ford (1951, p. 273) has described the use of liquid nuclear emulsion to study radioactive minerals in thin section. This highresolution method should be ideal when rocks and minerals of very low radioactivity are being examined, as the alpha tracks can be followed directly to their source.

We believe that the stripping-film method, described above, may be used without modification for histological studies and with some modification for polished-section and metallographic studies.

ACKNOWLEDGMENTS

This work was done as part of the program that the U. S. Geological Survey is conducting on behalf of the U. S. Atomic Energy Commission. We wish to acknowledge the assistance of Dr. Herman Yagoda, National Institute of Health, U. S. Public Health Service, who reviewed the manuscript and made valuable suggestions throughout the work. We are indebted to F. S. Reed and John Mergner of the Geological Survey for preparing thin sections.

References

- BERRIMAN, R. W., HERZ, R. H., AND STEVENS, G. W. W. (1950), A new photographic material—a high resolution emulsion for autoradiography: British Jour. Radiology, 23, 472–477.
- BOWIE, S. H. U. (1951), Autoradiographic techniques in geological research: Great Britain Geol. Survey, Bull., 3, 58-71.
- BOYD, G. A., AND WILLIAMS, A. I. (1948), Stripping film technique for histological autoradiographs: Proc. Soc. Exper. Biol. and Medicine, 69, 225-232.
- COPPENS, R. (1950), Sur l'emploi de l'émulsion photographique pour la détermination de la radioactivité des roches par l'examen des trajectoires des rayons α : Jour. Physique et radium, 11, 21-32.
- DONIACH, I., AND PELC, S. R. (1950), Autoradiographic technique: British Jour. Radiology, 23, 184–192.
- FORD, I. H. (1951), Radioactivity of rocks: an improvement in the photographic technique: *Nature*, 167, 273-274.
- GOODMAN, CLARK, AND THOMPSON, G. A. (1943), Autoradiography of minerals: Am. Mineral., 28, 456–467.
- HELLER, D. A. (1950), The radioautographic technique, AEC Document UCRL-579, 1-29.
- MACDONALD, A. M., COBB, JOCK, SOLOMON, A. K., AND STEINBERG, D. (1949), Stripping film technic for radioautographs: Proc. Soc. Exper. Biol. and Medicine, 72, 117-121.
- Mügge, O. (1909), Radioaktivität und pleochroitische Höfe: Centralbl. Mineralogie, 1909, pp. 114, 142.
- PELC, S. R. (1947), Autoradiography technique: Nature, 160, 749-750.
- PICCIOTTO, E. E. (1949), L'étude de la radioactivité des roches par la méthode photographique: Soc. belge géologie, Bull. 58, 75-90.
- POOLE, J. H. J., AND BREMNER, J. W. (1949), Investigation of the distribution of the main and a state of the photographic method: *Nature*, 163, 130-131.
- TYLER, S. A., AND MARAIS, J. J. (1941), A method for the determination of the relative radioactivity of mineral grains: *Jour. Sedimentary Petrology*, 11, 145–147.
- WEBB, J. H. (1948), Photographic plates for use in nuclear physics: Phys. Rev., 74, 511-532.
- WIENER, MARTIN, AND YAGODA, HERMAN (1950), Background eradication in thicklayered nuclear emulsions: Rev. Sci. Instruments, 21, 39-41.
- YAGODA, HERMAN (1946a), The localization of uranium and thorium minerals in polished section: Am. Mineral., 31, 87-124.

---- (1946b), Radiocolloid aggregates in uranium minerals: Am. Mineral., 31, 462-470.

- ------ (1949), Radioactivity measurements with nuclear emulsions, New York, John Wiley & Sons.
 - —, AND KAPLAN, NATHAN (1949), Spontaneous neutron emission from uranium and samarium: *Phys. Rev.*, **76**, 702–703.

Manuscript received July 21, 1951