



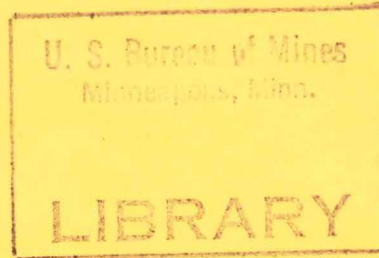
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**ASSESSMENT OF RESEARCH AND DEVELOPMENT NEEDS
AND PRIORITIES FOR ACID MINE DRAINAGE ABATEMENT**

(BCR Report L-822)

Prepared for

**UNITED STATES DEPARTMENT OF THE INTERIOR
BUREAU OF MINES**



By

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"The views and conclusions contained in this document are those of the authors and should not be interpreted as necessarily representing the official policies or recommendations of the Interior Department's Bureau of Mines or of the U. S. Government."

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16. Abstract In this report, recommendations are made for programs to develop workable techniques to stop or prevent acid mine drainage formation in those underground coal mines where acid production could take place. These recommendations include development of mine closure technology, development of surface water infiltration control procedures, study of the effect of roof collapse on mine water characteristics, evaluation of the long-term effects of flooding underground mines above drainage, investigation of the effectiveness of section sealing in active mines, development of a simple model for predicting mine water quality and quantity, preparation of a manual on mine drainage abatement procedures, and development of a program to disseminate information on mine drainage abatement to the coal industry. These recommendations are based on an in-depth evaluation of information acquired over years of research on acid mine drainage formation, on numerous laboratory and field studies of methods of preventing or minimizing acid mine drainage formation, and on review of the geological, hydrological, and mineralogical factors involved. There are 772 references.					
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FOREWORD

This report was prepared by Bituminous Coal Research, Inc., 350 Hochberg Road, Monroeville, Pa. 15146, under USBM Contract Number J0265044. The contract was initiated under the Coal Mine Health and Safety Program. It was administered under the technical direction of TCMRC, with Mr. William H. Engelmann acting as the Technical Project Officer. Mr. David L. Vila was the contract administrator for the Bureau of Mines.

This report is a summary of the work recently completed as part of this contract during the period June 30, 1976 to January 31, 1977. This report was submitted by the authors on June 27, 1977.

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I. CONCLUSIONS AND RECOMMENDATIONS

To accomplish the objectives of this project, nearly 800 documents, listed in the bibliography, Section V, were reviewed and evaluated. The index to the bibliography includes references to the following subject areas which were investigated: chemical and biological aspects of mine-drainage formation; abatement methods, including operating procedures, mine sealing, hydrologic controls, and use of various kinds of inhibitors; geologic parameters such as the conditions existing during coal formation, particularly those that affected pyrite content; water quality, including data on quality of ground water, streams and mine effluents, and information on coal characteristics.

In addition, researchers involved in mine-drainage control studies were contacted as were coal industry representatives who are responsible for mine-drainage control programs at active underground mines. BCR is indebted to these many experts who were willing to express candid opinions concerning past and present research as well as the need for development of new technology.

A significant percentage of the underground coal reserves may produce acid mine water when mined. During active mining, the quantity of acid mine water formed might be reduced, but not eliminated, by implementation of abatement measures that have been demonstrated previously. Any acid water that is formed can be neutralized before discharge to prevent surface water pollution. Other than in Appalachia, essentially all of the new underground mines will be situated below drainage; and, unless pumping is required to protect adjacent mines, will become flooded and have no discharges when abandoned. Flooding will eliminate the oxygen-pyrite contact, with the result that little additional acid will be formed. Even in the steeply rolling terrain that typifies Appalachia, most of the higher elevation coal seams have been mined and the majority of future mines will be located below drainage. However, many mines will still be situated above drainage, and procedures are not available for eliminating the acid discharge from them when abandoned. The major thrust of future Bureau of Mines' research, therefore, should be focused on developing methods of mine closure.

The specific conclusions and recommendations that have been drawn are as follows:

<u>Conclusions</u>	<u>Recommendations</u>
1. Although the exact mechanisms involved in acid formation by pyrite oxidation are not completely understood, the theories of AMD formation proposed by researchers at Ohio State University appear to most closely describe the chemical oxidation of pyrite in the natural mine environment.	1. Further research by the Bureau of Mines on acid mine drainage formation is not recommended. It would be interesting and, from a scientific viewpoint, should be continued, but it will not contribute to "at-source" abatement technology.

Conclusions

2. Sulfur and iron-oxidizing bacteria have been identified in natural mine drainages and have a role in acid mine drainage production. Although research has been carried on for a number of years by different investigators, there are still questions about the actual mechanism of bacterial action.
3. The occurrence of acid mine water cannot be correlated with the sulfur content or the abundance of pyrite in the coal. Other factors, including pyrite reactivity and the acid-forming potential or neutralization capability of adjacent rock structures, control the rate of acid formation.
4. Laboratory investigations, at best, poorly simulate the acid-formation reactions that take place at the mining site.
5. Mathematical modeling to predict flow and acid load has been successful for relatively small and simple mines. Even in these situations, too much field data are required for too many parameters.

Recommendations

2. Further research by the Bureau of Mines on biological processes in pyrite oxidation is not recommended. While it would be of academic interest and perhaps contribute to a better understanding of the role of bacteria in mine drainage formation, it would not advance "at-source" abatement technology.
3. Further research should be conducted to develop a method for determining the acid-forming potential of coal and adjacent stratigraphic measures that make up the natural mine environment.
4. A laboratory procedure should be developed for producing acid under all of the conditions which exist in the underground mine. This procedure should accurately simulate such variables as oxygen concentration, character of infiltrating water, moisture content, and product removal processes. Such a procedure is necessary to bridge the gap between the laboratory and the actual mine-drainage situation that exists in the field.
5. In order to make the modeling technique a useful and usable tool for predicting mine-water quality, a simple model, requiring the input of only a few critical variables, should be developed.

Conclusions

6. The most important factors influencing the quality and quantity of mine drainage are:
(a) pyrite presence, mode of occurrence (reactivity), and availability; (b) oxygen level; (c) bacteria; (d) quantity of water entering the mine; (e) quality of water entering the mine; and (f) water-flow pattern within the mine.
7. The basic mine drainage abatement principles are: (a) placement of acid-forming materials to eliminate or minimize contact with mine water; (b) inundation of acid-forming materials to eliminate contact with air; and (c) careful handling of water that does gain entrance to the mine to minimize contact with acid-forming materials and to expedite the water flow from the mine.
8. Each of five separate mining regions in the United States can be characterized by relative similarities in mine-water quality.
9. Acid mine drainage occurs mainly in the Northern Appalachian Region which includes Pennsylvania, Ohio, Maryland, and Northern West Virginia. However, even in this region, a large number of mines have drainages which are not acid.
10. Acid formation may be more widespread in the Illinois Basin than previously expected. However, the alkaline nature of the infiltrating water and the minimum amount of water produced at the mining sites, in general, precludes the acid formation

Recommendations

Conclusions

situation from developing into an acid mine drainage pollution problem.

11. While occasional acid production has been recorded for a number of coal seams, and while a few seams have been identified as having a probability of producing acid, no single seam has been shown to produce acid drainage at all times and at all places.
12. Based on surveys of the coal industry conducted by EPA and BCR, and on review of the mine drainage publications, 38 coal seams have been shown to produce acid mine drainage. However, in many instances, only a single reference could be found indicating that the seam produced acid.
13. In surface mining, mine drainage abatement procedures, in addition to reclamation technology, are sufficiently developed to minimize the problem when the mine is operating and to eliminate it when the mine is abandoned.
14. Underground mine-sealing technology is not sufficiently developed to guarantee that the mine-drainage problem will be eliminated when the mine is closed.
15. During mine or panel development, mine-water quality is similar to the quality of the local ground water.
16. Rock dusting has a positive effect on the quality of some mine drainages.

Recommendations

12. When opening a new mine in one of these coal seams, the coal industry should be alert to the possibility that acid problems may develop even though adjacent mines in the same seam have alkaline discharge.
13. Surface mining should be encouraged wherever possible for those coal seams located above drainage.
14. The thrust of the Bureau of Mines future research efforts should focus on developing mine-closure (sealing) technology.
16. Studies should be carried out to determine the extent to which rock dusting affects drainage quality.

Conclusions

17. The application of both biocides and bacteriophages in preventing acid mine drainage in actual underground mining environments is virtually an impossible task.
18. A number of chemicals have been investigated for their use either in coating pyritic surfaces or in reacting with these surfaces in order to render them inert and unable to initiate the formation of acid. While several substances have looked promising in laboratory investigations, chemical methods of inhibiting pyrite oxidation were not successful in mining environments.
19. Water infiltration control techniques can be used to reduce the volume of water that enters an underground mine.
20. Acid mine drainage abatement by interception of ground water before it enters the mine is in the theoretical stage of development and does not now appear practical.
21. The most effective abatement procedures that can be utilized in an operating mine involve minimizing the contact time between the water and acid-forming materials and removing the water as expeditiously as possible.
22. Development of a mine to the dip may greatly reduce the acidity of the mine discharge.

Recommendations

17. Further attempts to develop biological inhibition of pyrite oxidation are not recommended.
18. Although chemical inhibition of pyrite oxidation has academic interest, further attempts to develop underground "at-source" abatement techniques based on this method are not recommended. Some justification for future research in this area would be for treating coal-cleaning refuse piles. Indirectly, this would have a positive effect on underground mine-water quality, as such refuse could then safely be placed on the surface.
19. Further work on infiltration control should be carried out with appropriate monitoring to provide proper evaluation.
20. The results of pilot dewatering projects, currently being conducted, should be critically evaluated before additional research on dewatering techniques is considered.
21. A manual on Control of Acid Mine Drainage should be provided and, in order to achieve wider use of these practices, a program to disseminate this information throughout the coal industry should be developed.
22. Studies should be conducted to provide better methods of estimating the thickness of the barrier pillar or outcrop that will be required to maximize coal recovery and minimize barrier failure.

Conclusions

23. Caving that normally attends longwall mining could possibly reduce or inhibit acid mine drainage formation.
24. The maximum demonstrated effectiveness of air sealing in reducing acid loads is only about 50 percent.
25. Mine flooding appears to be the best method of minimizing or eliminating acid discharges from underground mines following abandonment. However, there are several drawbacks:
- (a) The tremendous volumes of water which will be contained may be inadvertently released with catastrophic results.
 - (b) The quality of the overburden ground water may be affected at those sites where a large hydrostatic head results from mining.
 - (c) A considerable amount of coal will be wasted when leaving outcrop and pillar barriers.
 - (d) Most mine-sealing projects were conducted at abandoned mines and the long-term effectiveness of sealing was not documented.
 - (e) Section sealing of abandoned portions of active mines may minimize mine drainage problems.

Recommendations

23. The effect of total roof collapse on mine drainage quality should be studied to determine whether roof collapse is a potential abatement method.
24. Further research on air sealing appears impractical and is not recommended.
- 25.
- (a) The Bureau of Mines should accept the responsibility of mapping these mines and monitoring all surface activities that might interfere with the integrity of the system.
 - (b) Future Bureau of Mines research should determine whether or not ground-water pollution might result from sealing.
 - (c) Further research should be conducted to determine pillar size requirements for providing adequate support for surface water bodies.
 - (d) The Bureau of Mines should sponsor a project to provide continuous monitoring, maintenance, and evaluation of some completed projects in order to determine the true effectiveness of mine sealing.
 - (e) Investigations into the effectiveness of section sealing should be conducted.

II. INTRODUCTION

A. JUSTIFICATION

Acid mine drainage is a naturally-occurring phenomenon. Where pyritic materials are exposed, they can be oxidized and the resulting oxidation products can be removed by water flowing across the reaction site. Such water will be acid in nature, and surfaces contacted by this water may be subject to a characteristic orange or "yellow boy" coating. Early records note that coal was discovered by observers who were aware that such seepages were often a sign of the presence of coal.

With the growth of the coal industry and the exposure of vast quantities of pyritic materials in many mines, acid mine drainage pollution increasingly became a problem. Although industrial and governmental efforts have been directed at varying rates since World War I at defining and eliminating this condition, acid mine drainage is still a serious effect of coal mining in many places. The work carried on over these past 60 years is well documented. It includes: research on the factors contributing to mine drainage formation, programs to control acid formation by chemical and biological means, implementation of mining practices to minimize the problem, and development of mine-drainage treatment processes.

As a result of these activities, methods have been developed that will virtually eliminate acid mine drainage pollution caused by surface and auger mining. Such methods include both "at-source" abatement procedures and mine-drainage treatment processes. The "at-source" abatement procedures involve the application of the two basic mine-drainage control principles; i.e., controlled placement of pyritic materials and efficient water handling. By careful implementation of these procedures, acid mine drainage formation and its subsequent discharge are greatly reduced.

According to the 1969 Appalachian Regional Commission Report, "Acid Mine Drainage in Appalachia," 19 percent of the acid load carried by Appalachian streams originates in active underground mines. The acid mine drainage control options available to operators of underground mines also include "at-source" abatement procedures and treatment. However, due to the nature of underground mining, "at-source" abatement procedures are exceedingly difficult, if not impossible, to implement. This load is being eliminated or greatly minimized by construction of mine-drainage treatment plants by the coal industry. However, construction and operation of such plants is costly, and treatment can only be considered a temporary solution to the problem. Temporary, in the sense that while stream pollution has been reduced, the source of the acid production has not been eliminated, and the stream may revert to an acid condition once mining has ceased. Temporary, also, since strikes, power failures, normal treatment-plant malfunctions, etc., may result in the occasional release of acid water in sufficient quantity to cause periodic pollution of the receiving stream.

Due to the increasing demand for coal, it is anticipated that underground mining will be greatly expanded in the near future. In order to provide permanent and reasonable cost control of acid mine drainage from underground coal mines, it is imperative that "at-source" abatement procedures be developed.

B. OBJECTIVES AND SCOPE

The purpose of this project is to recommend specific objectives of a research program that will lead to the development of "at-source" acid abatement procedures for active underground mines. This objective was approached by using available information: to make a critical evaluation of the "at-source" abatement work that has already been done; to describe the present knowledge concerning the formation of acid mine drainage and identify areas where more research is needed to more fully understand the problem; and to delineate the present and probable future extent of the acid mine drainage problem, and to characterize the environmental and mining parameters that contribute to its formation.

C. SUMMARY

The overall reaction which describes acid mine drainage formation is well documented in the literature. Acid mine drainage is formed as a result of the removal by mine water of the products formed by the oxidation of pyritic materials associated with the coal seam and the surrounding strata. Ferrous sulfate and sulfuric acid are the initial products of oxidation. Ferrous sulfate can subsequently be oxidized to ferric sulfate, and ferric ions can oxidize additional pyrite. Although numerous investigations have been carried out to define and to quantify the individual steps of the reaction and to determine which step is rate-controlling, a lack of information still exists concerning these basic aspects of acid formation.

The factors which contribute to the rate at which acid is formed are well defined and include the availability of oxygen and moisture. Increased acid production rates have also been attributed to the activity of sulfur-oxidizing and iron-oxidizing bacteria, which have been isolated consistently from acid mine drainage. Based on laboratory experiments to resolve the biological and chemical components of acid formation, investigators have attributed as much as a 17-fold increase in acid production directly to bacterial activity. The bacterial role in increasing the ferric-ferrous ratio in effluent water is well established. Microorganisms also have a significant influence on pyrite oxidation.

Another factor which controls the rate of acid formation is pyrite particle size. Laboratory studies have demonstrated that fine-grained pyrite was oxidized at a faster rate than coarser pyritic material. Other studies have indicated that the occurrence of acid mine drainage can be directly related to the occurrence of fine-grained (framboidal) pyrite.

The occurrence of acid mine drainage cannot be directly related to the sulfur content or the pyrite content of the coal being mined. According to

a recent survey, coal seams containing less than 1 percent sulfur rarely produce acid mine drainage, while those containing greater than 1 percent sulfur may or may not produce acid. It is apparent that other variables--including pyrite reactivity, oxygen concentration, character of infiltrating water, moisture content, and product removal processes--control the rate of acid formation and the opportunities for its release from the mine.

Mathematical models utilizing these variables, in addition to geologic and meteorologic data, to predict flow and acid load have been only moderately successful when applied to small, existing underground mines for which extensive field data were available. The difficulties of predicting acid formation from more complex or from proposed mines may be insurmountable.

Two approaches were used in evaluating the present and probable future extent of acid mine drainage occurrence. One method of assessing the problem was to compile and analyze available data on mine discharge quality. Information from two recent surveys was organized for each of five mining regions having generally characteristic water problems. The data were tabulated according to pH range of the mine effluent: less than four; from four through six; greater than six. In evaluating the data, all discharges having a pH below six were considered acid. This analysis showed that acid drainage was not a major problem for the coal industry taken as a whole. Even in the Northern Appalachian Region where acid drainage was most prevalent, approximately half or more of the mines surveyed reported the pH of their drainage to be above six.

In the second method of defining the problem, coal seams were evaluated for characteristics of the drainage they produced, and an attempt was made to identify acid-producing seams. Several coal seams in the Northern Appalachian Region were most likely to produce acid, but these seams were not invariably acid producers. A number of mines showed both acid and alkaline discharges. Water-handling techniques as well as mine hydrology and geochemistry can affect the character of mine discharges.

"At-source" abatement measures have been proposed based on eliminating one or more of the fundamental constituents of acid formation--bacteria, pyrite, water, and oxygen. "At-source" abatement also includes those procedures designed to minimize the transportation of acid from the reaction site or prevent its release from the mine.

Inhibition of acid-producing bacteria has been demonstrated in laboratory tests, but attempts to abate acid formation by applying inhibitors directly to underground mines were not successful. In the opinion of the authors the feasibility of successfully utilizing biological control to prevent acid formation in underground mines is very low for the following reasons; (1) The production and use of biocides may be prohibitive, both as to cost and toxicity; (2) The actual points of acid formation may not be accessible or perhaps not even known; (3) All of the infiltrating water would have to be inoculated; (4) Periodic applications would be necessary; (5) Acid formation from pyrite oxidation is a slow, continuing process occurring over long periods of time, and any inhibitor or coating applied for the purpose of abatement must be kept in place for equally long time periods; (6) Chemicals which have been

proposed to inhibit bacterial acid formation may result in additional pollutants which would have to be removed from the effluent; and (7) The chemical oxidation and leaching process would continue regardless of bacterial inhibition.

Coating of pyritic surfaces with an impervious shield to eliminate air and moisture contact with pyrite has been suggested as an "at-source" abatement control technique, but reported experience in this area is limited. The inaccessibility of acid-forming materials and the fact that new pyritic surfaces are constantly being exposed as a result of subsidence, make this control method impractical. Periodic applications would be necessary, and the additional pollutants that may result from any inhibitor would compound the problem.

If water could be completely eliminated from underground mines, acid mine drainage would not be formed. Although it is not possible to completely prevent water from entering underground workings, effective handling of water once it gains entrance to the mine can minimize water-pyrite contact. Careful mine planning can be used to minimize the mine's total production of water. Barriers can be left in place to reduce water infiltration through highly permeable or fractured overburden. A mine can be developed in such a manner as to take advantage of basic hydrological principles. For instance, mining down-dip, so that the mine can be completely flooded following closure, might eliminate all mine-water discharges. Diversion of surface waters away from mine openings is another way of reducing water infiltration into underground mines, thereby decreasing the potential pollution load. This technique has been successfully demonstrated in a number of projects. Careful placement of refuse would also reduce the amount of pollutants that may enter the effluent stream.

Longwall mining may be an effective and economical way to inhibit acid formation. The caving which results upon completion of mining may effectively bury acid-forming pyrites under a blanket of non-reactive materials, thereby reducing or eliminating air-pyrite contact.

Underground mine sealing to completely exclude air (air sealing) was attempted as early as the 1930's under Civil Works Administration (CWA) and Works Progress Administration (WPA) projects. The effectiveness of the air-sealing work accomplished during that period is inconclusive. In the 60's and 70's, projects to demonstrate air sealing were conducted in Pennsylvania and West Virginia. Evaluation of these programs has led to the conclusion that air sealing, at best, reduces acidity by only 50 percent. Periodic maintenance of the seals is necessary, and new air passageways are opened as a result of fracturing and subsidence. With every change in barometric pressure, the mine "breathes" thus replenishing its oxygen supply and negating any benefit derived from air sealing.

Mine flooding has been demonstrated to be the only viable method of minimizing or eliminating acid discharges from underground mines following abandonment. Abatement by flooding the mine is based on the fact that covering pyritic surfaces with water greatly decreases the amount of oxygen which can come in contact with pyrite. Reductions in acidity of as much as 99 percent have been reported when this technique was implemented.

Effective abandonment procedures that would eliminate post-abandonment acid mine drainage problems should be emphasized in future Bureau research programs. This research should include determination of width of barriers to be left in place and investigations into the effectiveness of section sealing of abandoned portions of active mines.

Since underground mining has historically been conducted in the Appalachian region, material relating to the acid mine drainage problem is most extensive for this region. Numerous data have been generated pertaining to sulfur in coal, geology of the associated strata, and stream and effluent quality. Many recent studies, particularly in Pennsylvania, have been conducted to evaluate the acid mine drainage problems of specific mining circumstances and to recommend abatement work that would eliminate these problems. Many of these abatement programs, utilizing mine-sealing and water-infiltration control techniques to demonstrate mine drainage abatement, have been completed. However, an evaluation of the long-term results of the abatement work is available for only a few projects. A monitoring program to evaluate the true effectiveness of these abatement projects should be undertaken, since valuable information can be gained from these experiences.

D. INVENTIONS

There are no patentable results or inventions from the work performed under this contract.

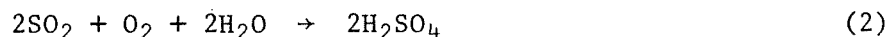
III. STATUS OF CURRENT KNOWLEDGE AND TECHNOLOGY

A. ACID FORMATION

1. Chemical Oxidation

a. Early Investigations: There is general agreement in the literature that the fundamental chemical reactions resulting in the formation of acid mine drainage involve the oxidation of pyritic sulfur in the presence of atmospheric oxygen and water to form sulfuric acid and iron sulfate. However, prior to the 1930's, few efforts were made to determine the mechanism of these reactions. During this early period considerable research was conducted on iron sulfides but not in relation to acid mine drainage.

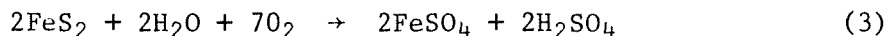
Using laboratory-controlled conditions to simulate the exposed coal seam, Burke and Downs^{148*} studied the effect of variables such as hydrogen ion, oxygen, and ferrous ion concentrations on the rate of pyrite oxidation. Based on their laboratory experiments, they proposed the following reaction sequence for pyrite oxidation:



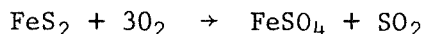
They postulated that since reaction (1) would proceed much slower than reaction (2), the formation of SO_2 was the rate-determining step in the oxidation of pyritic sulfur. It also was noted that sulfur balls reacted about 10 times faster than "museum grade" pyrite.

Nelson, *et al.*⁴⁹⁶ demonstrated that ferric sulfate assisted in the oxidation of pyritic sulfur and that oxygen content was an unimportant factor in the oxidation process. They also determined experimentally that pyritic sulfur in the finer sizes was oxidized at a faster rate than the larger sizes.

Early laboratory investigations at Ohio State University⁴⁹¹ involved basic research on the formation of acid in coal mines. In order to determine which mechanism was operative in acid formation, studies were made on iron disulfides. The Ohio State group found that oxygen was a critical factor in the rate of oxidation and that moisture influenced the rate. Since there was no indication that SO_2 was produced, they concluded that the pertinent reaction relating to acid mine drainage formation was



and not

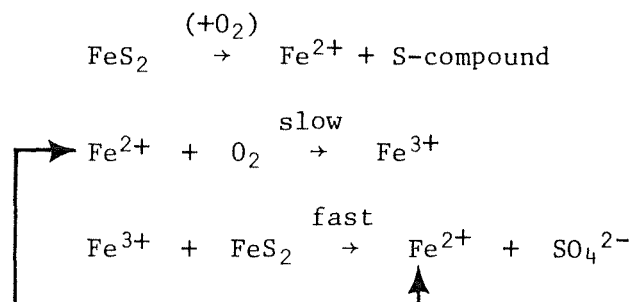


* Superscript numbers refer to items in the list of references, Section V.

as was stated by Burke and Downs. They also determined that the surface area of pyritic materials had a direct effect on acid formation.

The action of ferric sulfate solution on the rate of oxidation of pyrite was studied by Garrels and Thompson²⁹⁴ who postulated that between pH 0 and 2, the rate of oxidation was proportional to the fraction of pyrite surface occupied by ferric ion. They noted that the three pyrite specimens used, which were obtained from the Harvard Museum collection, oxidized at markedly different rates although the mechanism was the same.

b. Recent Studies: More recently, Stumm and others^{584,612,759} at Harvard University studied the rates of reactions related to mine-drainage formation. They proposed that FeS_2 reacted initially by a simple dissolution or by oxidation. They observed that in the absence of ferric iron no oxidation of iron pyrite occurred. Hence, the major oxidant of iron pyrite was ferric ion. Oxygen was involved only indirectly. It served to regenerate Fe^{3+} , which was the specific oxidant of pyrite. The oxidation rate was independent of pyrite surface area. The Harvard group concluded that the rate-determining factor in the oxidation of iron pyrite and subsequent formation of acid mine drainage was the oxidation of ferrous iron. Pyrite oxidation, as discussed by Singer and Stumm, can be represented as follows:



The Harvard group based their conclusions on data obtained from "museum grade" pyrite. Since this form of pyrite has been shown to be less reactive than naturally-occurring pyritic material found in coal measures, considerable doubt must be cast on the experimental results reported by the Harvard group and by other investigators who used "museum grade" pyrite.

Based on thermodynamic grounds, Clark^{174,176-177} refuted the theory by Singer and Stumm that pyrite went into solution by an ordinary chemical reaction. Since pyrite has a low solubility in water (1.48×10^{-2} mg/l at pH 2, 1.48×10^{-4} at pH 4, and 2.09×10^{-7} mg/l at pH 7), Clark postulated that the initial reaction resulting in pyrite going into solution was an electrochemical reaction; that is, a chemical oxidation-reduction reaction in which oxidation and reduction occurred at different locations on the pyrite surface. On thermodynamic grounds, both O_2 and Fe^{3+} can serve as oxidizing agents for pyrite.

Smith, Shumate, Sasmojo, and others^{490,559,590,591} at Ohio State University investigated the sulfide-to-sulfate reaction relative to acid mine drainage chemistry. They measured the oxidation rate of pyrite under aerobic and anaerobic conditions and demonstrated that oxidation involving either oxygen or ferric ion can be a major contributor to acid formation and that oxidation rates were dependent on pyrite surface area. However, they disagreed with Singer and Stumm's theory that the rate-determining step was the oxidation of ferrous iron.

The Ohio State group stated that for the mechanism proposed by Singer and Stumm to be rate-controlling, two conditions must be satisfied. First of all, pyrite oxidation by ferric ion should be at least equal to the oxidation of pyrite by oxygen. And secondly, the oxidation by oxygen of Fe^{2+} to Fe^{3+} in solution must be 14 times the rate of pyrite oxidation by O_2 . Under the experimental conditions used, the Fe^{3+} concentration had to be at least five times the Fe^{2+} concentration to satisfy the first condition. They determined that the aerobic oxidation rate of pyrite in a solution containing 0.01 gm-moles Fe was 5.0×10^{-6} gm-moles/hr-gm at pH 2.2. The rate of Fe^{2+} oxidation in a solution of 0.01 gm-moles/l Fe^{2+} was reported⁷⁷² to be 10^{-7} gm-moles/hr. These measurements yielded a ratio of pyrite oxidation to ferrous iron oxidation of only 0.02. Therefore, the second condition could not be satisfied. Based on these experiments, they concluded that the dissolved oxygen concentration was rate controlling in pyrite oxidation.

Smith, Shumate, et al. concluded that ferric ions may be an oxidizing agent for both aerobic and anaerobic oxidation. Based on experimental data, they proposed the following mechanism for the ferric iron oxidation of pyrite:

1. Ferric iron is adsorbed on a "reactive site" of pyrite to form an activated complex.
2. Ferrous ion is adsorbed on the "reactive site."
3. The activated complex decomposes to yield an adsorbed ferrous ion and transfer of electron.
4. Desorption of adsorbed ferrous ion occurs.

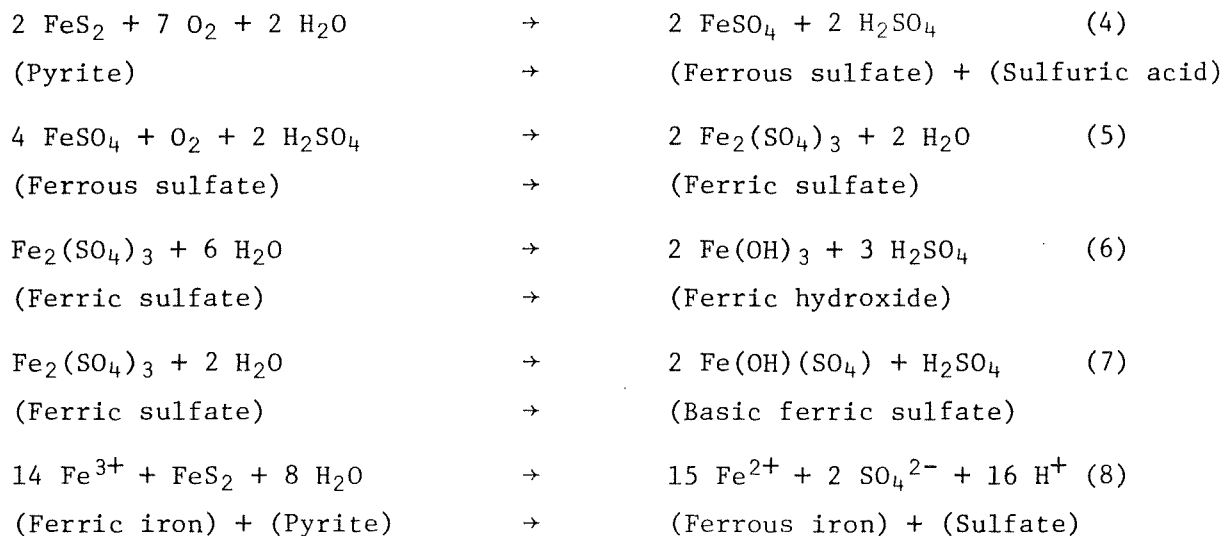
They proposed that Step 3, the surface transfer of electrons, was the rate-controlling step.

The Ohio State group⁵ also described the physical parameters affecting the rate of acid formation. They pointed out that pyrite is contained in a more-or-less porous matrix through which the oxygen must diffuse. A major rate-limiting factor in natural pyritic systems is, therefore, the transport of oxygen to the reaction site. Since air-saturated water does not contain enough dissolved oxygen to produce a significant acid load, and the rate of oxygen diffusion through a water-saturated strata is too low to be significant, they reasoned that the movement of oxygen to a reaction site must be through

a vapor phase; i.e., only that pyrite exposed to a vapor phase that is continuous to air will be oxidized at a significant rate.

Based on this conceptual model, the characteristics of the environment surrounding a reaction site in natural systems were described as a vapor phase saturated with water vapor producing a thin film (or filled pores) of water at the surface of the pyrite. The water in contact with the reaction sites would be essentially saturated with oxidation products.

c. Discussion: The following reactions, which describe the chemical oxidation of pyrite and subsequent reactions which may occur, are frequently reported in the literature:



Smith and associates at Ohio State University postulated that the rate-controlling step for acid formation in natural systems was the oxidation of pyrite by either oxygen or ferric iron as shown by reactions (4) and (8), respectively. Many workers demonstrated that the rate of pyrite oxidation was directly related to surface area^{133,491,496} and reactions (4) and (8) are dependent on this factor. Oxygen concentration was demonstrated to be a critical factor in pyrite oxidation.^{176,491,590} However, investigators at Harvard University dismissed direct oxidation by oxygen as insignificant. According to Stumm and associates, the rate-controlling step in acid formation was ferrous-iron oxidation as shown by reaction (5). Other workers postulated that reaction (5) occurred in solution remote from the pyritic surfaces and was not a rate-limiting reaction in acid formation. Reaction (5) was dependent on the number of bacterial cells and on the ferric- to ferrous-iron ratio.

The theories of Smith *et al.*, and Stumm *et al.*, evolved as the two major theories of AMD formation in the literature. The rate-controlling reaction proposed by Stumm was not consistent with findings such as the critical roles of oxygen concentration and of pyrite surface area in the rate of acid formation. These conflicts were explained by the fact that "museum grade" pyrite,

which was used by the Harvard group, was less reactive than pyrite associated with coal. The theories of Smith and Shumate were consistent with the findings of other investigators. Therefore, it must be concluded that the theories of AMD formation proposed by the Ohio State Group more closely describe the chemical oxidation of pyrite in the natural mine environment.

The exact mechanisms by which acid mine drainage is formed are not completely understood, but the gaps in the knowledge are not sufficient enough to warrant further extensive investigation. Further research in this area would be interesting, and from a scientific viewpoint should be continued, but it will not contribute to "at-source" abatement technology.

2. Microorganisms in Acid Mine Drainage: Work conducted at West Virginia University^{188, 357-363, 622-624, 626-628, 631, 634-635} during the 1940's and 1950's showed that bacterial action was involved in oxidation of ferrous iron to ferric iron in coal mine waters. An iron-oxidizing bacterium¹⁸⁷ and another bacterium similar to Thiobacillus thiooxidans were isolated from acid mine water.

Temple and Koehler⁶³² studied acid formation by Thiobacillus thiooxidans and Thiobacillus (T.) ferrooxidans in the laboratory. Acid formation was enhanced by the action of T. thiooxidans on sulfur, sulfur ball, and marcasite. No acid was formed with bacteria on pyrite. T. ferrooxidans oxidized ferrous iron formed from pyrite and increased the acid formation from sulfur ball and pyrite.

About the same time, Braley, Leathen, and associates^{136, 404, 406, 407} at Mellon Institute studied the effect of bacteria on pyritic materials. They also isolated Thiobacillus thiooxidans and iron-oxidizing bacteria from mine waters. Laboratory investigations demonstrated that T. thiooxidans increased acid production from marcasite and sulfur but not from pyrite or sulfur ball. The iron-oxidizing bacteria was shown to have an augmentative effect on acid formation from sulfur ball and marcasite but not from pyrite. The iron-oxidizing bacteria also oxidized ferrous sulfate. Leathen and Braley⁷⁴⁸ completely described the iron-oxidizing bacterium and proposed the generic name Ferrobacillus (F.) ferrooxidans.

A third group of investigators at Brigham Young University¹⁴⁷ in Utah isolated, characterized, and determined the nutrient requirements of two chemosynthetic autotrophic bacteria found in effluent streams. Both ferrous iron and sulfur were used as oxidizable substrates in the isolation of the bacteria. The culture grown on sulfur failed to oxidize ferrous iron; however, the culture isolated on ferrous iron was able to oxidize pyrite and sulfur. Since the bacteria corresponded most closely to the genus Thiobacillus, it was assumed that they were T. thiooxidans and T. ferrooxidans as reported by Leathen et al.⁴⁰⁸ and Temple and Colmer.⁶²⁸

A table summarizing the activities of the organisms studied by the three groups (Mellon Institute, West Virginia University, and the Utah group) was presented by Smith et al.³³¹ and is shown as Table 1 of this report.

TABLE 1. ACTIVITIES OF VARIOUS ORGANISMS ISOLATED FROM ACID MINE WATERS

<u>Substrates</u>	<u>T. thiooxidans</u>			<u>F. ferrooxidans</u>	<u>T. ferrooxidans</u>	
	<u>Mellon Group</u>	<u>W. Va. Group</u>	<u>Utah Group</u>	<u>Mellon Group</u>	<u>W. Va. Group</u>	<u>Utah Group</u>
Thiosulfate		+		-	+*	
Sulfur	+	+	+	(+; Silverman 1959)		+
Fe ⁺⁺	-	-	-	+	+	+
Sulfur Ball	-	+		+	+	
Pyrite	-	-	-	-	+	+
Marcasite	+	+		+		
Chalcopyrite			-			+
Molybdenite			-			+

+ = activity shown

- = no activity

Blank = not done

*Later, lost activity

Silverman et al.^{579,580} found that a significant oxidation of sulfur occurred by the bacterial action of F. ferrooxidans which were supplied to him by Leathen. Temple et al.⁶³² noted a difference in reactivities of different sulfur balls with Thiobacillus thiooxidans. He explained the disagreement with the results reported by Leathen in terms of a difference in the type of sulfuritic material used. Temple also found that Thiobacillus ferrooxidans enhanced acid formation from pyrite when the particle size was small enough. Since Leathen used coarse pyrite (between 1.0 and 2.4 millimeters or 0.04 and 0.095 of an inch) and Temple used a ball mill-grind pyrite, the reported results of the activity of F. ferrooxidans on pyrite may not be in disagreement.

A bacterium capable of oxidizing both sulfur and ferrous iron, Ferrobacillus sulfooxidans, was reported by Kinsel.⁷⁴⁴ However, since Silverman showed that F. ferrooxidans oxidized sulfur, the difference between the two bacteria was not apparent.

Walsh⁶⁷² reported the isolation from acidic iron-bearing streams of a stalked iron bacterium which belongs to the genus Metallogenium. The new bacterium catalyzed iron oxidation only in the pH range 3.5 to 5.0. Since Thiobacillus ferrooxidans were shown to catalyze ferrous iron oxidation below pH 3.5, Walsh hypothesized that stalked iron bacteria may be one of the major causes of the initial production of acidity and of the lowering of the pH in mine water.

Bacteria have been isolated consistently from acid mine drainage. Their role in acid formation will be discussed in the following section.

3. Chemical Versus Biological Acid Formation

a. Laboratory Studies: That bacteria are an integral part of the coal mine environment has been well established. The exact mechanism by which these organisms aid in acid formation is inconclusive. Various investigators have attempted to resolve the chemical and biological components of acid formation.

Leathen⁴⁰⁴ used substrates of sulfuritic materials which were prepared aseptically. The role of sulfur- and iron-oxidizing bacteria in the formation of acid was determined by examining inoculated and uninoculated substrates. He determined that Thiobacillus thiooxidans did not enhance the acid formation of sulfur ball or pyrite. A considerable increase in the amount of acid formed was noted in samples inoculated with iron-oxidizing bacteria as compared to uninoculated samples. A five-fold increase in acidity was reported with "sulfur ball" material, a four-fold increase with marcasite, and no increase with pyrite.

The influence of bacteria on acid formation was studied by Ashmead.⁷³ Iron disulfide and demineralized water were placed in a flask; the flask and contents were autoclaved. Natural mine water and sterile mine water were added to iron-disulfide suspension in order to ascertain the bacteriological extent of acid formation. Natural mine water, which was filtered through a bacterial filter, was also tested. Ashmead concluded that for every ton of sulfuric acid produced by chemical means, approximately four tons were produced by bacteriological means.

A Percolation Leaching Process was employed at Ohio State University⁴⁹¹ to resolve the acid-forming process into its fundamental components. The sulfuritic materials used were pyrite and marcasite. Samples were inoculated with acid mine water and sterile autoclaved water. The samples were then leached with tap water which was neutralized to pH of 7. The difference in acid production between inoculated and sterile samples was assumed to be indicative of biological activity. Both daily and cumulative acidities were

recorded for a period of 22 days. A small amount of acid was produced initially in both inoculated and sterile samples. Then there was a lag time, after which the inoculated samples produced about twice as much acid as the sterile samples. Fifty percent of the acid formed was attributed to bacterial activity. Marcasite produced approximately 17 times more acid than did pyrite. No special precautions were taken to maintain "sterile" conditions after the initial inoculations.

The standard Warburg manometric method was used by Silverman⁵⁸⁰ to study the oxidation of pyrites in the presence and absence of bacteria. Neither T. thiooxidans nor F. ferrooxidans enhanced the oxidation of crystalline pyrite. Both organisms accelerated the oxidation of marcasite, but T. thiooxidans did so only to a slight extent. The highest oxidation rates occurred with sulfur ball. Silverman demonstrated that sulfur-ball material which was inoculated with F. ferrooxidans underwent significantly greater oxidation than did sulfur-ball material with no bacteria present.

Smirnov⁵⁸⁶ attempted to determine whether the chemical or biological theory of acid formation was correct. He used natural, sterile, and disinfected mine waters and inoculated sterile water for his experiments. The type of sterile water and the method used for sterilization were not stated; however, sterile waters had significantly higher initial acidities than the natural waters used.

A flow of water was maintained through crushed pyrite for a period of 30 days. Smirnov reported that Thiobacillus thiooxidans and Thiobacillus ferrooxidans increased the rate of oxidation of pyrite to sulfuric acid by 11 to 13 times compared with oxidation under sterile conditions. He noted that when furacillin or formaldehyde was added to the water to achieve complete sterility, no acid was formed. Again, the acidities of these waters were high initially. Any increase, therefore, would have been small compared to the original acidity of the water.

Shearer et al.⁵⁷¹ were not specifically trying to resolve the chemical and biochemical theories of acid formation. However, their experiments resulted in some indication of how bacteria affect acid production. Separate coal piles were inoculated with F. ferrooxidans and with F. sulfooxidans. An uninoculated coal pile was used as a control. The coal piles were continuously leached with tap water over a six-month period. A significant increase in acid production from the inoculated piles compared to that from the uninoculated pile was reported.

Caruccio¹⁵⁸ conducted leaching studies to evaluate the acid potential of coal and coal-related materials. Unweathered coal and rock samples were taken from active and abandoned mines. The samples were ground to 1-2 mm in size and placed in leaching chambers. A continuous flow of humidified air was maintained over the samples. Twice a week the chambers were flushed and the acidities of the effluents were determined. Twenty days after the acid-generation trends were established, three samples were inoculated with bacteria from acid mine drainage pools. The acid-generation rate of one sample

increased six-fold; the other two samples showed no increase in the rate of acid formation.

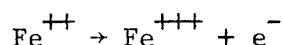
Silverman et al.⁵⁸¹ tested the effect of bacteria on coals of different rank. Distilled water acidified to pH 2.6 was used to leach bituminous, subbituminous, and lignite coal samples. Experiments were conducted in the presence and absence of F. ferrooxidans. All bituminous coals tested showed a considerable increase in pyrite oxidation in inoculated compared to uninoculated samples. The pyrites associated with subbituminous coals and lignites seemed to resist bacterial attack.

A study of the oxidizing characteristics of pyritic materials from various coal mines was conducted by Lorenz and Tarpley.⁴²⁶ The effect of F. ferrooxidans was determined by using inoculated and sterile samples. The type of drainage from the mine where each pyrite sample was collected was noted. Of the eleven pyritic materials tested, eight samples showed at least a 17-fold increase in oxidation due to bacterial activity. Four of the eight samples were associated with alkaline drainage areas. Two samples, one from an acid-drainage area and the other from a section which had both acid and alkaline waters, were not affected by bacterial activity.

Work conducted by Lazaroff⁴⁰² suggested that the growth of T. ferrooxidans required a high proportion of sulfate ions to other anions. Schnaitman et al.⁵⁶⁰ corroborated this finding.

Walker and Randles⁶⁵⁸ postulated that iron bacteria catalyze the initial removal of electrons from the iron of the pyrite surface, thereby initiating the acid-formation process. They pointed out that the microorganisms that were associated with acid mine drainage formation grew only in the presence of oxygen, although they did not dismiss the idea that other oxidizing agents may substitute for oxygen.

Lundgren and Schnaitman⁴³⁴ suggested that iron-oxidizing bacteria catalyze the oxidation of ferrous iron by atmospheric oxygen by using the energy released by the reaction:



They postulated that a rate-limiting step in pyrite oxidation may be ferric ion formation.

Duncan, Landesman and Walden²⁵⁷ proposed that ferric ion was of minor importance in the oxidation of sulfide minerals and demonstrated that direct bacterial action on iron and sulfide was the mode by which acid was produced.

Baker and Wilshire⁸⁰⁻⁸³ demonstrated that F. ferrooxidans and T. thiooxidans accelerated oxidation of ferrous iron and S_2^{2-} but did not alter the pyritic dissolution rate.

Stumm-Zollinger⁶¹⁴ suggested that the rate of oxidation appeared to be proportional to the quantity of bacteria present.

b. Discussion: Large increases in pyritic oxidation rates have been observed in microbially catalyzed reactions in the laboratory. However, the environment at the pyritic-reaction site in these laboratory oxidations is not representative of conditions in natural pyritic systems. An artificially high ferric-ferrous ratio is generated in a large quantity of water solution that is in contact with a relatively small surface area of exposed pyrite. In natural pyritic systems, pyrite in contact with this much water would be "flooded," with negligible oxygen in the vicinity of the pyrite surface.

It has been shown experimentally that both low water-to-pyrite ratios and a high concentration of acid and salts in solution reduce the number and activity of bacteria to the point that the ferric ion oxidation of pyrite is insignificant compared to oxygenation. Both of these conditions pertain to reaction sites in underground mines where vapor phase oxidation must be involved.

Ample evidence shows the existence of bacteria and their effectiveness in increasing the ferric-ferrous ratio in effluent water. However, the question remains as to whether microorganisms have a significant influence on the rate of acid production in natural pyritic systems. The key question is where do the microbially catalyzed reactions occur; in ponded water after contacting pyrite, or at the reaction site. If in ponded water, there would be no significant change in acid production due to the presence of bacteria simply because there is no further contact between the water and the pyrite. If bacteria are sufficiently active at reaction sites to generate a ferric-ferrous ratio greater than 5 to 1, then bacteria have a significant effect on acid production. In view of the nature of the environment at the reaction site, as determined in many previous Ohio State University studies, it does not seem likely that bacteria are capable of generating a ferric-ferrous ratio approaching 5 to 1.

Although effluent water from a pyritic system may have a high ferric-ferrous ratio, this does not imply a similar situation at reaction sites since effluent water is not representative of the water in contact with reaction sites.

Mathematical simulation models of pyritic systems such as that suggested by Morth⁴⁸⁶ indicated the acid production observed in the field could be explained solely by the oxygenation rate of pyritic materials. Microbially-enhanced reaction rates were not needed to account for the rate at which acid was produced. It should be pointed out that utilization of the model requires physical information such as the porosity of the pyrite-containing strata and chemical data including pyrite reactivity. Good methods for determining these parameters are not available.

The bacterial role in acid formation is not well defined. Further research to determine the relative importance of biological processes in

pyrite oxidation would contribute to a better understanding of acid formation but would not advance "at-source" abatement technology. Even if abatement measures based on inhibiting bacteria could be developed, chemical oxidation would still continue and acid would still be formed. In addition, the application of an effective bacterial inhibitor would present insurmountable problems.

4. Pyrite Reactivity: Laboratory studies demonstrated that pyrite in finer sizes was oxidized at a faster rate than coarser pyritic material. Braley¹¹⁹ determined that marcasite reacted ten times faster than yellow pyrite and that black amorphous pyrite reacted eighty times faster than yellow pyrite.

X-ray diffraction analyses were made on field samples collected from coal mines in Ohio, Pennsylvania, and West Virginia.⁴²⁶ The coal seams represented were the Pittsburgh, Lower Freeport, and Lower Kittanning seams. The only iron sulfide identified was pyrite. Marcasite was not found in any of the samples. This observation has been confirmed by many investigators.

Caruccio⁷²³ indicated that the occurrence of acid mine drainage can be directly related to the occurrence of framboidal pyrite within a coal seam and associated strata. Framboidal pyrite was described as a fine-grained, highly-reactive form of pyrite. High percentages of framboidal pyrite were associated with the marine-brackish water coals and a paucity of framboidal pyrite was related statistically to fresh water coals.

Caruccio¹⁵⁸ compared two areas of strip mining: one area underlain by rocks of the "marine-brackish water," Lower and Middle Kittanning formations; and the other area underlain by the "fresh water" Freeport strata. He found that the occurrence of acid mine drainage could not be related to the sulfur content of the coals since one area produced acid waters and the other area produced non-acid discharges, but the sulfur content of the coals was similar. A microscopic examination of the coal samples revealed that in coals from the acid-producing area, a major portion of the pyrite occurred as clusters of spheres approximately 25 microns in diameter, while in samples from the non-acid-producing area, the pyrite commonly had a massive form with grains greater than 400 microns.

According to Renton and Hidalgo,⁵⁴⁴ "there is no correlation between the amount of acid produced and the abundance of pyrite in coal." The only quarrel with this statement is that it presumes that a minimum level of pyrite is always present. As can be seen by Tables 2 and 3, coal seams with low (less than 1 percent) sulfur values, and correspondingly lower pyrite values, rarely produce acid mine drainage. These tables are based on a 1972 BCR survey and include data from both surface and underground mines. Table 2 shows the mine-drainage quality of coal seams in the Northern Appalachian region and the average sulfur content of the mines working the seam. Table 3 shows, on a nationwide basis, mine-drainage quality by mine compared to the average sulfur content of the coal being mined. These data tend to corroborate Caruccio's conclusion that the occurrence of acid mine drainage cannot be related to the sulfur content of the coal.

TABLE 2. RELATIONSHIP OF SULFUR IN COAL TO MINE WATER QUALITY
(by Coal Seams in Northern Appalachia)

Coal Seams	Coal Sulfur Content in Percent					Total Mines
	< 1.0	1.1-2.0	2.1-3.0	3.1-4.0	> 4.1	
Lower Kittanning						
pH < 4.0	--	1	1	4	--	6
pH 4-6	--	--	--	--	1	1
pH > 6.0	--	--	--	--	--	--
Middle Kittanning						
pH < 4.0	--	--	2	--	--	2
pH 4-6	--	--	1	--	2	3
pH > 6.0	--	1	2	1	1	5
Lower Freeport						
pH < 4.0	--	1	1	--	--	2
pH 4-6	--	--	--	--	--	--
pH > 6.0	--	--	4	2	--	6
Upper Freeport						
pH < 4.0	--	2	--	--	--	2
pH 4-6	--	--	1	--	--	1
pH > 6.0	--	4	4	--	--	8
Pittsburgh						
pH < 4.0	--	3	4	8	2	17
pH 4-6	--	2	--	--	--	2
pH > 6.0	--	3	5	6	5	19
Meigs Creek						
pH < 4.0	--	--	--	--	--	--
pH 4-6	--	--	--	--	--	--
pH > 6.0	--	--	--	2	--	2
Total						
pH < 4.0	--	7	8	12	2	29
pH 4-6	--	2	2	--	3	7
pH > 6.0	--	8	15	11	6	40

TABLE 3. RELATIONSHIP OF SULFUR IN COAL TO MINE WATER QUALITY

<u>Geographical Regions</u>	<u>Coal Sulfur Content in Percent</u>					<u>Total Mines</u>
	<u>< 1.0</u>	<u>1.1-2.0</u>	<u>2.1-3.0</u>	<u>3.1-4.0</u>	<u>> 4.1</u>	
Northern Appalachia						
pH < 4.0	--	7	8	12	2	29
pH 4-6	--	2	2	--	3	7
pH > 6.0	--	8	16	11	6	41
Southern Appalachia						
pH < 4.0	2	2	1	--	--	5
pH 4-6	10	4	--	--	--	14
pH > 6.0	70	12	--	--	--	82
Illinois Basin						
pH < 4.0	--	--	--	2	2	4
pH 4-6	--	--	--	3	2	5
pH > 6.0	--	2	7	12	7	28
Western Interior Province						
pH < 4.0	--	--	--	--	--	--
pH 4-6	--	--	--	--	1	1
pH > 6.0	1	--	--	3	3	7
Western Provinces						
pH < 4.0	--	--	--	--	--	--
pH 4-6	--	--	--	--	--	--
pH > 6.0	7	--	--	--	--	7
Total						
pH < 4.0	2	9	9	14	4	38
pH 4-6	10	6	2	3	6	27
pH > 6.0	78	22	23	26	16	165

It can be concluded from these data that coal seams containing less than 1 percent sulfur rarely produce acid mine drainage. On the other hand, those coals containing greater than 1 percent sulfur, and therefore probably containing a significant amount of pyrite, may or may not produce acid mine water. It cannot be concluded that a seam containing 4 percent sulfur will produce acid drainage on more occasions, or will have greater acidity, than a seam containing only 1 to 2 percent sulfur. It is generally accepted that only the inorganic fraction (normally pyrite) of the sulfur present in coal contributes to acid formation. In addition, it is apparent that other factors, including pyrite reactivity, control the rate of acid formation.

The reactivity of pyrite has been experimentally measured and the effect of mineralogical properties on the reactivity semi-quantitatively described. However, a method for determining the reactivity of the pyrite associated with coal and adjacent rock measures that make up the natural mine environment is not available. A quantitative determination of the acid-producing potential of a particular coal or shale strata should be developed and would aid in predicting the probability and extent of acid mine drainage in a specific area.

There is also a need to develop, for use in the laboratory, a procedure for producing acid under all of the various conditions which exist in the underground mine. This procedure must accurately simulate such variables as oxygen concentration, character of infiltrating water, moisture content, and product-removal processes. Although numerous past laboratory studies have been based on the utilization of such a procedure, a reliable method does not exist. The development of this procedure would bridge the gap between laboratory and field work and may lead to a better understanding of the chemical processes involved in acid formation in the natural mine environment.

5. Predicting Mine-water Quality: Numerous factors have been shown to contribute to the quality and quantity of the mine-water discharge. Since the underground mine environment is a dynamic heterogeneous system, making predictions relative to the characteristics of the mine drainage to be formed is a complex task. Mathematical models may be the answer, and several have been developed to predict water discharge quantity and acid load from existing underground mines. These models, if perfected, are expected to accurately predict the environmental response to imposed changes in order (a) to determine the long-term effects of proposed mining operations, (b) to evaluate the effectiveness of abatement measures, and (c) to plan optimum treatment facilities.

The Water Generation Model (WATGEN) developed at Pennsylvania State University⁵⁰⁹ required input parameters such as soil and rock properties, meteorologic data, fluid properties, and aquifer characteristics. Using these data, the water-discharge quantity from an active mine was simulated for one year. The simulated flows were generally higher than actual pumpages, but the predicted and the actual mine-water discharges did not differ by more than 22.5 percent.

The Acid Mine Drainage Quantity and Quality Generation Model developed at the Ohio State University⁵⁴⁷ was based on two models: a modified version of the Stanford Watershed Model (SWM), to calculate the amount of moisture reaching the mine aquifer; and the Acid Mine Drainage Model (AMD) described by Morth et al.,⁴⁹⁰ to simulate pyrite oxidation kinetics and oxidation product removal. The major inputs to the total model are: climatological, watershed characteristics, and mine characteristics data. The inputs are represented by 64 parameters as listed below. The outputs from the total model include: average daily mine water discharge, the associated acid concentration or load, plus the average daily flow in the receiving streams or at basin outlets. In using the Ohio State Model to simulate acid loads and water quantity from the McDaniel's Mine, the actual and predicted values were in fair agreement.

LIST OF INPUT PARAMETERS FOR OHIO STATE UNIVERSITY ACID MINE
DRAINAGE QUANTITY AND QUALITY GENERATION MODEL

SWM Parameters

Measurable Parameters

Impervious area that drains directly into the stream channel
Watershed drainage area in square miles
Index capacity of the existing channel in cubic feet per second
Empirical constant for convection
Estimate of the stream and lake surface area as a fraction of AREA
Daily interflow recession constant
Long term ratio of average basin rainfall to average watershed ppt.
Daily baseflow recession constant
Streamflow routing parameter for low flows
Stream routing parameter for flood flows
Mean overland flow path length in feet
Average ground slope in feet/foot of the overland flow surfaces

Trial and Adjustment Parameters

Index controlling the rate of infiltration
Index to estimate interception, depression storage capacity of the soil surface
Index to controlling time distribution, quantities of moisture entering interflow
Index for estimating soil surface moisture storage capacity
Factor relating infiltration rates to evaporation rates for seasonal adjustment
Minimum value of EN
Current value of groundwater slope index in inches
Daily baseflow recession adjustment factor
Current soil moisture storage, in inches
Soil profile moisture storage index, in inches
Groundwater storage increment, in inches

Assigned or Selected Parameters

- Maximum interception rate for dry watershed
- Soil evaporation parameter
- Index for groundwater flow leaving basin
- Groundwater evaporation parameter
- Manning's n for overland flow on soil area
- Manning's n for overland flow on impervious area
- Index for routing
- Current soil moisture storage

AMD Parameters

Mine System

- Watershed area of mine in square miles
- Number of air-solid waterface increments
- Number of layers in coal seam
- Number of depth increments in model
- Length of depth increments in feet
- Length of air-solid interface increments in feet
- Datum plane for top of coal seam in feet
- Literal description of stratum
- Elevation of stratum relative to datum plane in feet

Pyrite Oxidation Kinetics

- Oxygen consumption rate of pyrite
- Void fraction of the stratum
- Mine temperature correction factor
- Volume occupied by gram mole gas in cubic feet
- Constant for calculating rate constant

Oxidation Product Removal

- Aquifer storage in inches
- Constant relating mine water flow and aquifer storage
- Alkalinity conversion factor
- Minimum flow rate to cause acid removal by flooding
- Minimum flow rate to cause acid removal by leaching
- Constant to determine the inundated distance
- Hypothetical slope of the water level
- Fraction of stored products removed daily by inundation
- Base gravity diffusion constant

Basin and Mine Characteristics

Basin Characteristics

- Drainage area, sq miles
- Length of principal water course, feet

Average slope percent
Peak discharge of record, cfs
Land use

Mine Characteristics

Area, sq ft
Average height, ft
Peak minewater flow recorded, gallons per day
Peak acid load recorded, lb per day
Principal materials in coal seam

Thirty-three watersheds were evaluated¹⁰¹ to determine data availability to apply the SWM-AMD model. Data on acid-production parameters and mine conditions were generally lacking. These parameters, which are perhaps the most critical input data for accurate modeling, must be estimated.

Mathematical modeling was successful as far as predicting flow and acid load from several existing underground mines for which extensive field data were available. These mines were relatively small and well-defined compared to many underground mines, yet the input data were voluminous both in terms of the number of parameters and the number of determinations that had to be made to assign values for each of the parameters. In order to develop a more generally applicable model, a wide range of underground mine systems must be simulated and the results checked with field data.

The idea of a generalized model to predict acid formation from proposed and existing mines is intriguing, but the difficulties of modeling such complex and dynamic systems may be insurmountable. The completed model may be so complex that only the few who developed the model, and not coal-industry personnel in general, could actually apply it. The need, therefore, is to develop a simple model that would require input of only a few critical variables to estimate potential acid production.

B. SCOPE OF ACID MINE DRAINAGE PROBLEM

1. Data Base: A considerable volume of information and data was evaluated to determine the present and probable future extent of the acid mine drainage problem. Such information included:

- a. Water-quality information available in the publications in the BCR mine drainage library;
- b. Water-quality data for specific mine sites that were generated by the 1974-75 study by the Environmental Protection Agency to develop Effluent Limitation Guidelines for the coal industry;
- c. BCR file data relating mine-water quality to coal seams and to the specific sulfur content of the mines sampled;
- d. U.S. Bureau of Mines data on underground mining reserves;
- e. Reports on the geologic and hydrologic variables that influence mine drainage formation and discharge;
- f. Descriptions of various mining systems and their effect on mine drainage formation and discharge; and
- g. Discussion with mining experts.

In reviewing both the technical and non-technical literature, it became apparent that at present there exists a general misconception concerning the extent of the acid mine drainage problem. To most people, and this includes some who have worked on the problem for years, acid mine drainage is a normal result of mining, no matter what type of mining is undertaken or where the mining is located. In reality, the reports specifically referencing the occurrence of acid mine drainage and the data collected during BCR and Environmental Protection Agency (EPA) surveys of the coal industry, as presented in Tables 4 and 5, indicate that most drainage from coal mines is alkaline. Acid mine water is a problem that is limited to a relatively small percentage of coal seams and mines.

In 1972, BCR surveyed the coal industry to determine the extent of environmental problems on a mine-to-mine basis. Mine water quality data developed through the survey can be found in Tables 2 to 8 of this report. Most of these tables also include mine water quality information determined by Skelly and Loy Engineers, Harrisburg, Pennsylvania, as part of their 1975 study of the coal industry for EPA. The purpose of this study was to develop a data base to be used by EPA to establish Effluent Limitation Guidelines for the coal industry. It is noted that approximately 30 percent of the mines included in the EPA study were also covered by the BCR survey. In the tables, the mines are categorized on the basis of the worst (most acid) discharge from the mine. For example, if a mine had multiple discharges and only one was acid, the mine is included in the category reflecting the lowest pH. Three categories of pH

ranges < 4, 4-6, and > 6 are used, with all discharges having a pH less than 6 being considered acid. It is recognized that pH is, at best, only a partial indicator of mine-water quality. Nevertheless, it is the parameter which is normally available for most discharges and, therefore, the one used in describing the extent of the mine drainage problem.

The general mine-water quality, on a nationwide basis, of all mines included in the surveys is presented in Table 4. Of the 248 mines responding to the BCR survey, 65, or 26 percent of the mines reporting, produced acid water. In the EPA study, only 41 of 163 mines, or 25 percent of the mines for which data were available, had acid discharges or had discharges that required neutralization.

Included in the 248 mines responding to the BCR survey were 170 underground mines which represented more than 40 percent of the 1972 underground production. As shown in Table 5, only 49 of these 170 mines, or 29 percent of the active underground mines, produced acid discharges.

2. Regional Diversification of Acid Mine Drainage Formation: Obviously, numerous factors influence both the quality and the quantity of mine drainage that is produced at any given mining site. In all cases, these factors are variables which are dependent on either or both the geologic and geographic location of the mine and the total mining system employed. The most important factors, as discussed elsewhere, are: (a) pyrite presence, mode of occurrence (reactivity), and availability; (b) oxygen level; (c) bacteria; (d) quantity of water entering the mine; (e) quality of water entering the mine; and (f) water-flow pattern within the mine.

The coal industry's experience is that, geographically, the country can be divided into five separate mining regions and that the mines within each region display similar mine-water quality problems. These are: Region 1--Northern Appalachia which includes Pennsylvania, Ohio, Maryland, and Northern West Virginia; Region 2--Southern Appalachia which includes Southern West Virginia, Eastern Kentucky, Virginia, North Carolina, Tennessee, and Alabama; Region 3--Illinois Basin which includes Indiana, Illinois, Michigan, and Western Kentucky; Region 4--Western Interior Province which includes Iowa, Nebraska, Missouri, Kansas, Oklahoma, Arkansas, and Texas; and Region 5--the Western Provinces which include North Dakota, South Dakota, Montana, Idaho, Wyoming, Colorado, Utah, New Mexico, Arizona, Oregon, and Washington. This same regional breakdown is used in the following discussion.

TABLE 4. GENERAL NATIONWIDE MINE WATER QUALITY

<u>Data on Mine Water Quality</u>	<u>Number of Mines Reporting</u>	
	<u>BCR Survey</u>	<u>EPA Effluent Guideline Study</u>
Mine water quality pH < 4	37	31
Mine water quality pH 4-6	25	5
Mine water quality pH > 6	147	86
Dry or no data available	18	6
No mine water data but survey forms indicate:		
(a) neutralization provided	3	5
(b) no neutralization required	18	30
Total Mines in Data Base	248	163

TABLE 5. NATIONWIDE MINE WATER QUALITY FOR UNDERGROUND MINES

<u>Data on Mine Water Quality</u>	<u>Number of Mines Reporting</u>
Mine water quality pH < 4	31
Mine water quality pH 4-6	15
Mine water quality pH > 6	93
Dry or no data available	11
No water quality data but survey forms indicate:	
(a) neutralization provided	3
(b) no neutralization required	17
Total Underground Mines	170

With the exception of the Appalachian Basin, the literature that has specifically been devoted to identifying the extent of the acid mine drainage problem is limited. In the Appalachian Basin, particularly Northern Appalachia, the Appalachian Regional Commission, the U.S. Environmental Protection Agency, the USGS, and the states of Pennsylvania, Ohio, West Virginia, and Maryland have devoted considerable effort to delineating the effects of mine drainage.

The relative intensification of activity in Appalachia is due not only to the concentration of mining in the East but also to the limited existence of a general acid mine drainage problem in the other regions.⁷²⁴ As can be seen from the data in Table 6, presenting the results of the BCR and EPA surveys on the regional basis, the majority of mines with highly acid (pH < 4) discharges are located in Northern Appalachia.

Table 7 shows the same categories of data for the regions as Table 6 but includes underground mines only. It can be seen, by comparing these two sets of data, that acid mine drainage is generated by approximately the same percentage of underground mines as surface mines.

During discussions of this project with coal industry representatives and mine drainage experts, it was emphasized that the similarity ends there. In surface mining, mine drainage abatement including reclamation technology is sufficiently developed to minimize the problem when the mine is operating and to eliminate it when the mine is abandoned. In underground mining, abatement principles are well defined but are difficult or impossible to implement.

The basic principles are: (1) placement of acid-forming materials to eliminate or minimize contact with mine water; (2) inundation of acid-forming materials to eliminate contact with air; and (3) careful handling of water that does gain entrance to the mine to minimize contact with acid-forming materials and to expedite the water flow from the mine. A second reason why surface mining results in fewer mine-drainage problems is that underground mine sealing technology is not sufficiently developed to guarantee that the problem will be eliminated when the mine is abandoned.

It is recommended, therefore, that, wherever possible, surface mining should be encouraged for those coal seams located above drainage; and the thrust of the Bureau of Mines future research efforts in acid mine drainage abatement should focus on developing mine closure (sealing) technology.

TABLE 6. MINE EFFLUENT DATA BY REGIONS

Regions	Number of Mines Reporting						Total Mines in Survey	Percent of Mines Acid or Requiring Neutral- ization
	Mine Water Quality			Dry or No Data Avail- able	No Mine Water Data but Survey Forms Indicate:			
	pH < 4	pH 4-6	pH > 6		Neutral- ization Provided	No Neutral- ization Required		
Northern Appalachia								
BCR Survey	28	7	41	5	1	0	82	44
EPA Survey	24	5	25	0	5	6	65	52
Southern Appalachia								
BCR Survey	5	12	68	8	2	14	109	17
EPA Survey	2	0	25	2	0	10	39	5
Illinois Basin								
BCR Survey	4	5	24	2	0	4	39	23
EPA Survey	4	0	14	0	0	2	20	20
Western Interior Province								
BCR Survey	0	1	7	0	0	0	8	12
EPA Survey	1	0	5	2	0	3	11	9
Western Provinces								
BCR Survey	0	0	7	3	0	0	10	0
EPA Survey	0	0	17	2	0	9	28	0
Total								
BCR Survey	37	25	147	18	3	18	248	26
EPA Survey	31	5	86	6	5	30	163	26

TABLE 7. UNDERGROUND MINE EFFLUENT DATA (BCR SURVEY)

Regions	Number of Mines Reporting						Total Mines in Survey	Percent of Mines Acid or Requiring Neutral- ization
	Mine Water Quality			Dry or No Data Avail- able	No Mine Water Data but Survey Forms Indicate:			
	pH < 4	pH 4-6	pH > 6		Neutral- ization Provided	No Neutral- ization Required		
Northern Appalachia	26	4	29	3	1	0	63	49
Southern Appalachia	4	9	57	5	2	13	90	17
Illinois Basin	1	2	6	2	0	4	15	20
Western Interior Province	0	0	0	0	0	0	0	0
Western Provinces	0	0	1	1	0	0	2	0
Total	31	15	93	11	3	17	170	29

Another approach to specifically delineating potential acid-problem areas was to attempt to identify the acid-producing seams. Again, much more information was available for the two Appalachian regions than for any other region.

According to the Appalachian Regional Commission Report Acid Mine Drainage in Appalachia, "In Pennsylvania, Ohio, and northern West Virginia, the Monongahela Group is responsible for acid mine water pollution in varying degrees. The Sewickley and the Redstone seams produce a slight acid to acid discharge in some localities, while at others the water discharge may remain non-acid. The Pittsburgh seam sometimes produces acid water in underground mines. This varies from locality to locality, and the water discharge varies from non-acid to acid depending upon the mine location in the coal basin. The Allegheny Group, also, is responsible for acid mine water pollution in northern Appalachia. The Upper and Lower Freeport seams produce slight acid to acid water discharges in some localities, while at others the water may remain non-acid. The Upper, Middle, and Lower Kittanning seams produce slight acid to acid water discharges in many localities, while at others the water discharges may remain non-acid. The Lower Kittanning seam often discharges acid water from both strip and underground mines."

"In southern West Virginia, western Virginia, eastern Kentucky and Tennessee, and northern Alabama, the Pottsville Groups of bituminous coal seams produce occasional slight acid water discharges, though the mine drainage pollution problem in the southern part of Appalachia is less severe when compared with the same problem in the northern part of Appalachia. The Island Creek coal seam in Logan County, West Virginia occasionally produces slight acid water discharges. The Pocahontas coal seams apparently produce little or no acid water discharges. The coal seams in eastern Kentucky and Tennessee and in northern Alabama produce occasional acid water discharges in some localities, while in others the water discharge may remain non-acid."⁷

Other work in Appalachia included that conducted by the West Virginia Geologic Survey, who sampled and determined the relative acid-producing potential of 19 coal seams in West Virginia.^{544,545} The 19 coal samples represented the oldest through the youngest seams from the coal measures (Pennsylvanian and Dunkard strata). The greatest amount of acid was produced by the Pittsburgh and Upper Freeport coals, which exceeded the next highest acid produced, the Bakerstown coal, by a factor of 10. However, the acid-producing values of the Pittsburgh coal showed less variation than did those for the Upper Freeport, which indicates that estimating the relative acid-producing potential of the Upper Freeport coal would be more of a problem than estimating that of the Pittsburgh seam. Of all the coals tested, only the Pittsburgh, Upper Freeport, and Bakerstown were determined to be major acid producers. The Waynesburg coal yielded relatively low acid-production values, averaging five times less than those of the Bakerstown. Coal samples from the following seams showed extremely low acid-production values: Washington, Uniontown, Redstone, Elk Lick, Harlem, Brush Creek, Kittanning, Lower Mercer, Cedar Grove, No. 2 Gas, Sewell, Pocahontas No. 3, and Peacock.

These reports, as can be seen from Table 8, are in agreement with the data from the mines sampled during the BCR and EPA surveys.

TABLE 8. MINE-DRAINAGE QUALITY OF COAL SEAMS
IN NORTHERN APPALACHIA

Coal Seams	Number of Mines Reporting Effluent Water Quality					
	pH < 4		pH 4-6		pH > 6	
	BCR	EPA	BCR	EPA	BCR	EPA
Meigs Creek	0	0	0	0	2	0
Pittsburgh	17	11	2	3	19	9
Upper Freeport	2	4	1	0	8	8
Lower Freeport	2	1	0	0	6	3
Upper Kittanning	0	0	0	0	1	0
Middle Kittanning	2	3	3	2	5	5
Lower Kittanning	6	5	1	0	0	0

It is recognized and is demonstrated by the available data that there are significant differences in the mine-water quality produced by the various coal seams. Using the BCR survey, the EPA survey, and by searching the BCR mine drainage library, it was determined that, at least on occasion, the coal seams listed in Table 9 produce acid mine drainage.

It is important to note that, although these coal seams are listed as potential acid producers, in many instances only a single reference could be found indicating that the seam produced acid. In most cases, there are significant differences in the quality of water produced by mines in the same seam but at different locations. When opening a new mine in one of these seams, the industry anticipates that the water quality will be the same as adjacent mines in the same seam. Nevertheless, if the seam produces acid at other locations, the rule of thumb is to be alert to the possibility that acid problems might develop at any given site even though nearby mines may have alkaline discharges.

TABLE 9. LIST OF COAL SEAMS* THAT ON OCCASION HAVE BEEN KNOWN TO PRODUCE ACID MINE DRAINAGE

029	Meigs Creek, Sewickley	135	Chilton
033	Redstone	151	Cedar Grove, Elkhorn No. 3
036	Pittsburgh	154	Lower Cedar Grove, Up. Elkhorn No. 2
055	Barton	157	Alma, Blue Gem
063	L. Bakerstown	168	No. 2 Gas, Eagle No. 2 Gas
067	Bakerstown, Brush Cr.	174	Upper Eagle
070	Mahoning	176	Eagle
071	Upper Freeport	195	Hagy
073	Lower Freeport	207	Clements, Gwin
076	Upper Kittanning	212	Lily
077	Princess No. 7	285	Lower Seaboard
080	Middle Kittanning	294	Stearns
084	Lower Kittanning	334	Pocahontas No. 6
087	Clarion	484	Kentucky No. 11
090	Princess No. 5	489	Kentucky No. 9, Springfield No. 5
095	Brookville	492	Bevier
103	Stockton, Lewiston	502	Seelyville III
111	Coalburg	506	Davis
121	Winifrede	906	Rowe

*The three-digit numbers refer to the coal bed numbers used by the Bureau of Mines IC 8680 "The Reserve Base of U.S. Coals by Sulfur Content."

Differences in water quality also often occur within the same mine. For instance, 26 of the 63 mines included in the BCR survey of Northern Appalachia had multiple discharges. Sixteen of these mines are shown in the acid category (pH < 6) in Table 7, although at least 50 percent of the discharges from 11 of these 16 mines were alkaline. The general reasons given by industry experts for the differences in mine-water quality include:

- (1) During mine or panel development, the mine-water quality is usually alkaline and is similar to the quality of the local ground water;
- (2) The corollary to this is that during development the mine water may be acid, as sumps and effective water-handling procedures have not been established;
- (3) The effectiveness of abatement procedures, such as water handling, may differ within the mine with the result that some discharges are acid;

(4) In some instances it is suspected, but not documented, that rock dusting has a positive effect on the quality of some discharges;

(5) The quality of the water draining from abandoned areas of the mine will be influenced by the type of material that is present in the roof. As the roof collapses, this material becomes a significantly more important factor in affecting water quality than does the coal. If it contains reactive pyrites, an acid mine water will be formed. If limestone or calcareous shales are present, the water will be alkaline.

Due to the lack of complete data, it is impossible to reliably map those areas where acid mine drainage can be expected to be formed as a result of underground mining of bituminous coal. Mine-drainage maps that are presently available generally reference stream water quality and do not differentiate between the effects of surface mining and underground mining.

Tables 10-14 do provide some insight into the possible extent of the underground acid mine drainage problem on a state and county basis for each of the coal regions. Extreme caution is urged in using these data as they are based on several tenuous assumptions, and the resulting tables reflect a "worst case" situation. These assumptions include:

(1) Each coal seam included in Table 8 is a potential acid producer if the coal-bed sulfur content exceeds one percent. This ignores the effect of differences in pyrite reactivity which has been demonstrated to exist;

(2) Each numbered coal seam is considered a potential acid producer no matter where it exists, even if there is only a single reference available indicating it produces acid. This assumes that coal-bed characteristics are inter- and intra-regionally translatable;

(3) Local geographical and hydrologic conditions are the same throughout the country. This assumption ignores the fact that many mid-western and western mines are essentially dry or that in all regions influent ground water, or other materials available in the coal or roof rock, may contribute sufficient alkalinity to provide neutralization of any mildly-acid water that may be formed during mining.

Table 15 provides a summary of Tables 10-14 and indicates the percentage of the total underground bituminous coal reserve base that, assuming the worst-case situation, could have an acid mine drainage problem when mined. These percentages are understandably higher than those found in Table 6, which lists the percentage of mines found by ground surveys to have acid discharges. They do follow the same regional trends, however, in that Region 1 has the highest acid mine-water incidence and production potential, followed by Regions 3, 2, 4, and 5.

TABLE 10. REGION 1 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION
(million short tons)

State and County	Total Underground Bituminous Coal Reserves	Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming		
		< 1% Sulfur	Sulfur Value	
			Unknown	
Maryland				
Allegany	378.6	1.5	0.0	377.0
Garrett	523.3	89.5	0.0	429.4
Maryland Total	901.9	91.0	0.0	806.4
Ohio				
Athens	1,327.0	68.3	12.9	1,245.8
Belmont	3,927.1	0.0	76.0	3,513.2
Carroll	757.7	0.0	36.3	721.4
Columbiana	747.9	17.7	0.0	706.5
Coshocton	127.3	0.0	3.5	123.9
Gallia	339.6	0.0	10.7	328.9
Guernsey	1,184.4	0.0	517.3	667.1
Harrison	1,523.3	0.0	507.9	624.3
Hocking	205.7	1.9	9.3	196.3
Holmes	28.5	0.0	1.1	27.3
Jackson	154.7	0.0	16.2	138.4
Jefferson	1,356.2	3.3	0.0	455.7
Lawrence	477.0	5.0	0.0	472.0
Mahoning	308.2	0.0	0.0	308.2
Meigs	396.3	0.7	0.0	395.6
Monroe	468.4	0.0	468.4	0.0
Morgan	435.1	0.0	29.9	405.2
Muskingum	720.8	0.0	1.1	719.7
Noble	570.0	0.0	14.6	555.4
Perry	645.0	17.5	0.0	627.5
Scioto	4.8	0.0	1.3	3.5
Stark	376.6	0.0	5.5	371.1
Tuscarawas	841.2	0.0	21.3	813.0
Vinton	301.2	0.0	10.8	290.4
Washington	196.0	0.0	0.0	196.0
Wayne	3.3	0.0	2.9	0.4
Ohio Total	17,423.3	109.4	1,747.0	13,906.8

TABLE 10. REGION 1 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Continued)
(million short tons)

State and County	Total Underground Bituminous Coal Reserves	Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming		
		< 1% Sulfur	Sulfur Value Unknown	> 1% Sulfur
Pennsylvania				
Allegheny	881.4	27.8	455.0	398.6
Armstrong	1,092.9	16.7	62.7	1,013.5
Beaver	435.3	36.2	59.1	340.0
Bedford	89.5	2.5	0.2	86.8
Blair	18.7	1.9	1.6	15.2
Bradford	12.8	0.0	12.8	0.0
Butler	863.8	6.4	0.0	853.4
Cambria	1,454.2	272.4	0.0	1,181.8
Cameron	25.4	0.0	25.4	0.0
Centre	228.3	12.1	0.8	215.4
Clarion	640.1	11.7	69.9	558.5
Clearfield	1,102.5	93.5	0.0	1,008.9
Clinton	40.8	0.0	25.6	15.1
Elk	86.1	2.3	0.0	83.8
Fayette	1,023.7	19.6	0.0	814.6
Fulton	10.8	0.0	10.8	0.0
Greene	6,515.7	42.4	1,011.9	4,283.1
Huntingdon	36.3	1.9	0.0	34.3
Indiana	1,716.9	126.8	25.3	1,564.8
Jefferson	407.4	57.1	1.6	348.7
Lawrence	136.9	0.0	63.5	73.3
Lycoming	11.4	0.0	1.4	10.0
McKean	146.4	0.0	0.0	146.4
Mercer	128.0	0.7	86.4	40.9
Somerset	1,240.6	56.7	0.0	1,175.7
Tioga	36.5	2.2	0.0	20.9
Venango	55.0	0.0	0.4	54.5
Washington	3,604.2	143.8	187.2	2,904.2
Westmoreland	747.5	33.9	22.9	686.3
Pennsylvania Total	22,788.9	968.6	2,124.5	17,928.7

TABLE 10. REGION 1 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Concluded)
(million short tons)

State and County	Total Underground Bituminous Coal Reserves	Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming		
		Sulfur Value		
		< 1% Sulfur	Unknown	> 1% Sulfur
Northern West Virginia				
Barbour	948.8	0.0	43.0	398.7
Braxton	467.7	183.0	90.4	194.4
Brooke	169.3	0.0	38.1	131.1
Calhoun	0.3	0.0	0.0	0.0
Clay	695.0	460.1	135.2	99.7
Doddridge	248.4	0.0	239.2	0.0
Gilmer	154.1	0.0	0.0	154.1
Grant	313.2	81.6	0.0	231.6
Hancock	132.4	0.0	18.3	114.1
Harrison	380.9	0.0	0.0	380.6
Kanawha	1,120.6	238.9	0.0	199.7
Lewis	730.3	57.3	0.0	669.8
Marion	2,599.5	104.3	54.3	2,102.5
Marshall	3,043.8	0.0	0.0	2,840.7
Mason	116.7	0.0	0.0	115.4
Mineral	218.5	49.0	0.9	115.0
Monongalia	3,008.8	47.4	0.0	2,371.5
Ohio	379.0	0.0	79.9	291.0
Preston	837.5	34.8	2.9	799.8
Putnam	142.5	0.0	0.0	141.4
Roane	19.4	1.9	0.0	17.5
Taylor	388.2	0.0	0.0	388.2
Tyler	79.6	0.0	0.0	0.0
Upshur	876.9	42.3	112.5	224.9
Wayne	403.6	71.0	261.6	71.0
Wetzel	846.1	0.0	743.8	0.0
Northern West Virginia Total				
	18,321.1	1,371.6	1,820.1	12,052.7

TABLE 11. REGION 2 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION
(million short tons)

State and County	Total Underground Bituminous Coal Reserves	Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming		
		< 1% Sulfur	Sulfur Value Unknown	> 1% Sulfur
Alabama				
Blount	0.7	0.0	0.0	0.0
Cullman	28.1	0.0	0.0	0.0
DeKalb	1.3	0.0	0.0	0.0
Fayette	93.7	0.0	0.0	0.0
Jackson	13.2	0.0	0.0	0.0
Jefferson	757.9	0.0	0.0	66.0
Marion	118.4	0.0	0.0	0.0
St. Clair	1.9	0.0	0.0	0.0
Shelby	3.8	0.0	0.0	0.0
Tuscaloosa	143.7	0.0	0.0	0.0
Walker	630.0	0.0	0.0	0.0
Winston	5.3	0.0	0.0	0.0
Alabama Total	1,798.1	0.0	0.0	66.0
Eastern Kentucky				
Bell	106.4	26.0	40.0	3.0
Boyd	51.1	0.0	20.0	0.0
Breathitt	410.6	17.5	0.0	84.0
Carter	33.2	0.0	6.5	17.0
Clay	138.8	28.0	6.4	72.0
Clinton	0.5	0.0	0.5	0.0
Elliott	33.6	0.0	33.5	0.0
Floyd	952.6	489.0	77.4	369.0
Greenup	24.9	0.0	17.0	0.0
Harlan	1,408.5	722.0	133.0	299.0
Jackson	48.0	0.0	13.4	0.0
Johnson	78.0	11.5	24.0	35.0
Knott	1,248.4	531.0	146.0	218.0
Knox	50.0	21.0	0.0	5.0
Laurel	5.4	0.0	0.0	3.0

TABLE 11. REGION 2 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Continued)
(million short tons)

State and County	Total Underground Bituminous Coal Reserves	Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming		
		< 1% Sulfur	Sulfur Value Unknown	> 1% Sulfur
Eastern Kentucky (continued)				
Lawrence	56.5	111.0	18.0	0.0
Lee	25.2	0.0	1.6	16.3
Leslie	620.0	312.0	5.0	78.6
Letcher	730.4	318.0	0.0	247.0
McCreary	146.1	0.0	0.0	78.0
Magoffin	214.4	38.0	0.0	94.0
Martin	249.6	12.0	145.0	0.0
Menifee	3.7	0.0	3.7	0.0
Morgan	29.0	0.0	14.2	0.0
Owsley	2.0	0.0	2.0	0.0
Perry	560.2	8.0	21.4	40.1
Pike	2,170.0	1,341.1	65.7	380.4
Powell	0.8	0.0	0.8	0.0
Pulaski	28.2	0.0	21.4	0.0
Rockcastle	1.6	0.0	1.1	0.0
Wayne	1.6	0.0	1.6	0.0
Whitley	15.6	2.7	0.0	12.2
Wolfe	22.1	0.0	14.4	0.0
Eastern Kentucky Total	9,466.5	3,988.8	833.6	2,051.6
North Carolina				
Chatham	12.2	0.0	0.0	0.0
Lee	19.1	0.0	0.0	0.0
North Carolina Total	31.2	0.0	0.0	0.0

TABLE 11. REGION 2 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Continued)
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value</u>	
			<u>Unknown</u>	<u>> 1% Sulfur</u>
Tennessee				
Anderson	89.3	7.9	0.0	50.4
Bledsoe	15.1	0.0	0.0	0.0
Campbell	194.7	48.5	0.0	98.4
Claiborne	35.9	1.4	22.1	4.5
Cumberland	21.2	0.0	0.0	0.0
Fentress	39.8	0.0	0.0	0.0
Grundy	8.9	0.0	0.0	0.0
Hamilton	36.5	0.0	0.0	0.0
Marion	26.5	0.0	0.0	0.0
Morgan	56.7	0.2	0.0	18.5
Overton	4.2	0.0	0.0	0.0
Pickett	0.2	0.0	0.0	0.0
Putnam	17.8	0.0	0.0	0.0
Rhea	7.6	0.0	0.0	0.0
Roane	1.3	0.0	0.0	0.0
Scott	58.8	9.0	0.0	14.0
Sequatchie	44.7	0.0	0.0	0.0
Van Buren	0.2	0.0	0.0	0.0
White	7.8	0.0	0.0	0.0
Tennessee Total	667.1	67.0	22.1	185.8
Virginia				
Buchanan	820.0	73.9	0.0	194.1
Dickenson	515.5	0.8	0.0	2.5
Lee	167.4	31.4	0.0	136.3
Russell	403.0	0.0	0.0	0.0
Scott	44.6	0.0	0.0	0.0
Tazewell	166.0	3.9	0.0	16.0
Wise	716.6	101.2	10.4	32.8
Virginia Total	2,833.2	211.2	10.4	381.7

TABLE 11. REGION 2 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Concluded)
(million short tons)

State and County	Total Underground Bituminous Coal Reserves	Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming		
		< 1% Sulfur	Sulfur Value Unknown	> 1% Sulfur
Southern West				
Virginia				
Boone	1,868.8	712.6	0.0	845.1
Fayette	796.3	484.9	34.3	56.9
Greenbrier	269.7	77.3	0.0	45.1
Lincoln	360.4	113.4	118.0	129.0
Logan	3,076.3	1,121.9	0.0	1,501.8
McDowell	912.4	63.9	19.4	11.7
Mercer	38.6	5.6	0.5	2.2
Mingo	1,887.4	875.3	0.7	654.2
Nicholas	1,433.7	746.3	63.8	232.5
Pocahontas	131.6	73.4	0.0	0.0
Raleigh	1,656.3	328.7	96.5	252.4
Randolph	757.2	317.2	16.3	75.9
Summers	5.9	5.9	0.0	0.0
Tucker	121.0	34.2	0.0	86.7
Webster	1,098.9	242.2	129.8	355.4
Wyoming	1,642.0	274.6	78.8	77.2
Southern West				
Virginia Total	16,056.5	5,477.4	558.1	4,326.1

TABLE 12. REGION 3 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value Unknown</u>	<u>> 1% Sulfur</u>
Illinois				
Adams	0.02	0.0	0.0	0.0
Bond	1,831.4	0.0	79.4	1,752.1
Bureau	1,029.4	0.0	0.0	250.2
Cass	13.1	0.0	3.4	0.0
Christian	3,347.4	0.0	617.2	2,651.3
Clark	168.5	0.0	103.2	0.0
Clinton	1,322.5	0.0	2.1	1,320.0
Coles	80.7	0.0	44.0	0.0
Crawford	442.6	0.0	291.0	0.0
Douglas	411.7	0.0	11.0	390.6
Edgar	1,749.9	0.0	906.0	332.0
Edwards	54.0	0.0	54.0	0.0
Fayette	1,173.7	0.0	877.7	0.0
Franklin	3,038.4	802.0	168.0	1,905.0
Fulton	220.5	0.0	0.0	77.6
Gallatin	1,761.1	0.0	0.0	1,139.0
Greene	52.1	0.0	0.0	0.0
Grundy	246.0	0.0	0.0	32.5
Hamilton	2,440.2	0.0	2,440.0	0.0
Henry	28.3	0.0	0.0	2.6
Jackson	226.7	0.0	0.0	194.0
Jefferson	1,800.6	128.0	0.0	1,672.6
Jersey	42.0	0.0	8.3	0.0
Kankakee	79.9	0.0	0.0	0.0
Knox	68.0	0.0	16.6	0.0
LaSalle	1,083.0	0.0	0.0	92.0
Lawrence	893.6	0.0	527.7	0.0
Livingston	586.5	0.0	0.0	48.9
Logan	813.7	0.0	0.0	813.7
McClellan	421.0	0.0	0.0	107.0
Macon	439.3	0.0	18.9	420.4
Macoupin	3,421.1	0.0	56.6	3,007.0
Madison	1,366.5	0.0	0.0	1,214.0
Marion	421.0	0.0	102.0	319.0
Marshall	358.0	0.0	9.8	0.0

TABLE 12. REGION 3 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Continued)
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value Unknown</u>	<u>> 1% Sulfur</u>
Illinois				
(continued)				
Menard	1,460.0	0.0	0.0	1,460.0
Mercer	12.6	0.0	0.0	0.0
Montgomery	3,906.6	0.0	209.9	3,269.0
Morgan	144.8	0.0	0.0	43.4
Moultrie	123.1	0.0	0.0	123.1
Peoria	289.2	0.0	0.0	213.0
Perry	1,201.0	0.0	0.0	1,201.0
Putnam	588.9	0.0	78.9	0.0
Randolph	214.0	0.0	0.0	220.0
Rock Island	12.8	0.0	0.0	0.0
St. Clair	951.4	0.0	19.3	932.1
Saline	2,553.4	4.9	0.0	2,314.8
Sangamon	3,540.0	0.0	0.0	3,514.0
Scott	0.1	0.0	0.0	0.0
Shelby	712.5	0.0	534.1	53.1
Tazewell	69.3	0.0	0.0	34.2
Vermilion	1,544.3	0.0	44.5	537.0
Wabash	262.0	0.0	262.0	0.0
Washington	1,555.2	0.0	12.1	1,543.0
Wayne	89.0	0.0	89.0	0.0
White	992.5	0.0	628.6	360.0
Williamson	1,573.1	83.0	0.0	1,194.0
Woodford	213.9	0.0	42.3	0.0
Illinois Total	53,441.9	1,017.9	8,257.6	34,753.2

TABLE 12. REGION 3 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Continued)
(million short tons)

State and County	Total Underground Bituminous Coal Reserves	Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming		
		< 1% Sulfur	Sulfur Value Unknown	> 1% Sulfur
Indiana				
Clay	168.2	0.0	0.0	53.3
Daviess	109.3	0.0	18.7	0.0
Dubois	5.5	0.0	0.0	0.0
Fountain	7.0	0.0	0.0	0.0
Gibson	1,301.5	0.0	0.0	883.0
Greene	254.4	0.0	0.0	197.0
Knox	1,453.5	0.0	25.0	652.5
Martin	0.01	0.0	0.0	0.0
Parke	57.2	0.0	0.0	19.0
Perry	10.1	0.0	0.0	0.0
Pike	244.8	4.0	0.0	210.0
Posey	720.9	0.0	412.0	0.0
Sullivan	1,922.3	153.0	0.0	706.0
Vanderburgh	451.4	0.0	203.0	234.0
Vermillion	497.6	0.0	0.0	369.0
Vigo	1,212.1	0.0	0.0	721.0
Warrick	532.6	0.0	21.0	333.0
Indiana Total	8,948.5	157.0	679.7	4,377.8
Western Kentucky				
Butler	20.2	0.0	0.0	0.0
Christian	2.5	0.0	0.0	0.0
Crittendon	11.6	0.0	0.0	0.0
Daviess	126.2	0.0	70.9	48.8
Edmondson	0.1	0.0	0.0	0.0
Grayson	0.2	0.0	0.0	0.0
Hancock	0.5	0.0	0.0	0.0
Henderson	1,503.6	0.0	17.4	1,411.2
Hopkins	1,805.7	0.0	0.0	1,300.0
McClellan	723.6	0.0	96.4	627.2
Muhlenberg	898.4	0.0	0.0	711.9
Ohio	264.6	0.0	6.6	242.8
Union	1,926.3	0.0	0.0	1,807.7
Webster	1,436.4	0.0	15.9	1,216.2
Western Kentucky Total	8,719.9	0.0	207.2	7,365.8

TABLE 12. REGION 3 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Concluded)
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value Unknown</u>	<u>> 1% Sulfur</u>
Michigan				
Bay	55.6	0.0	0.0	0.0
Genesee	7.0	0.0	0.0	0.0
Huron	6.5	0.0	0.0	0.0
Saginaw	27.0	0.0	0.0	0.0
Shiawassee	1.8	0.0	0.0	0.0
Tuscola	19.7	0.0	0.0	0.0
Michigan Total	117.6	0.0	0.0	0.0

TABLE 13. REGION 4 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value Unknown</u>	<u>> 1% Sulfur</u>
Arkansas				
Crawford	19.1	0.0	0.0	0.0
Franklin	6.1	0.0	0.0	0.0
Scott	12.5	0.0	0.0	0.0
Sebastian	268.3	0.0	0.0	0.0
Arkansas Total	306.0	0.0	0.0	0.0
Iowa				
Appanoose	157.4	0.0	0.0	157.4
Boone	120.6	0.0	0.0	0.0
Cass	0.4	0.0	0.0	0.0
Dallas	50.4	0.0	0.0	0.9
Davis	37.6	0.0	0.0	0.0
Decatur	33.5	0.0	17.1	0.0
Greene	13.3	0.0	0.0	0.0
Guthrie	4.7	0.0	0.7	0.0
Hamilton	8.5	0.0	0.0	0.0
Hardin	4.6	0.0	0.0	0.0
Henry	1.2	0.0	0.0	0.0
Jasper	85.3	0.0	0.0	0.0
Jefferson	48.9	0.0	0.0	0.0
Keokuk	77.5	0.0	0.0	0.0
Lee	2.7	0.0	0.0	0.0
Lucas	134.4	0.0	1.9	0.0
Mahaska	325.3	0.0	0.0	0.0
Marion	398.3	0.0	0.0	0.0
Marshall	3.2	0.0	0.0	0.0
Monroe	518.7	0.0	0.0	0.0
Muscatine	0.1	0.0	0.0	0.0
Polk	492.7	0.0	0.0	0.0
Scott	2.2	0.0	0.0	0.0
Story	34.0	0.0	0.0	0.0
Van Buren	48.7	0.0	0.0	0.0
Wapello	148.9	0.0	0.0	0.0
Warren	87.5	0.0	1.0	0.0
Webster	43.9	0.0	0.0	0.0
Iowa Total	2,884.9	0.0	20.7	158.3

TABLE 13. REGION 4 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Continued)
(million short tons)

<u>State and County</u>	Total Underground Bituminous Coal Reserves	Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming		
		<u>< 1% Sulfur</u>	<u>Sulfur Value</u>	
			<u>Unknown</u>	<u>> 1% Sulfur</u>
Missouri				
Adair	482.2	0.0	0.0	44.0
Audrain	469.7	0.0	0.0	0.0
Barton	109.4	0.0	0.0	0.0
Bates	292.3	0.0	0.0	0.0
Boone	60.7	0.0	60.7	0.0
Caldwell	128.6	0.0	128.6	0.0
Callaway	163.9	0.0	163.9	0.0
Cass	72.9	0.0	0.0	0.0
Chariton	63.8	0.0	0.0	3.7
Clay	32.4	0.0	0.0	32.4
Daviess	74.0	0.0	0.0	74.0
Grundy	31.0	0.0	0.0	31.0
Harrison	223.7	0.0	0.0	0.0
Henry	130.2	0.0	0.0	0.0
Howard	152.9	0.0	152.9	0.0
Johnson	67.5	0.0	0.0	0.0
Linn	564.7	0.0	0.0	0.0
Macon	370.5	0.0	0.0	225.3
Mercer	589.3	0.0	0.0	0.0
Montgomery	141.3	0.0	0.0	0.0
Nodaway	17.3	0.0	0.0	0.0
Putnam	384.7	0.0	0.0	384.7
Randolph	670.4	0.0	0.0	670.4
Ray	109.9	0.0	0.0	109.9
St. Clair	209.8	0.0	0.0	0.0
Sullivan	373.6	0.0	90.3	0.0
Vernon	67.4	0.0	0.0	0.0
Worth	19.4	0.0	0.0	0.0
Missouri Total	6,073.6	0.0	596.4	1,575.4

TABLE 13. REGION 4 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Concluded)
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value</u>	
			<u>Unknown</u>	<u>> 1% Sulfur</u>
Oklahoma				
Atoka	3.3	0.0	0.0	0.0
Coal	36.8	0.0	0.0	0.0
Craig	156.3	0.0	0.0	0.0
Haskell	84.1	0.0	0.0	0.0
Latimer	63.0	0.0	0.0	0.0
LeFlore	269.7	0.0	0.0	0.0
Okfuskee	2.3	0.0	0.0	0.0
Okmulgee	106.2	0.0	0.0	0.0
Pittsburg	138.3	0.0	0.0	0.0
Oklahoma Total	860.1	0.0	0.0	0.0

TABLE 14. REGION 5 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value Unknown</u>	<u>> 1% Sulfur</u>
Colorado				
Archuleta	92.1	0.0	0.0	0.0
Delta	65.8	0.0	0.0	0.0
Fremont	180.3	0.0	0.0	0.0
Garfield	553.0	78.2	0.0	0.0
Gunnison	696.6	0.0	0.0	6.5
Huerfano	278.3	8.7	0.0	0.0
LaPlata	322.1	0.0	0.0	0.0
Las Animas	832.0	0.0	13.7	0.0
Mesa	229.4	0.0	16.3	0.0
Moffat	1,994.8	0.0	202.1	0.0
Montezuma	19.1	0.0	0.0	0.0
Montrose	42.8	0.0	17.1	0.0
Park	25.3	0.0	0.0	0.0
Pitkins	88.6	0.0	0.0	0.0
Rio Blanco	1,067.4	0.0	175.4	0.0
Routt	2,738.7	0.0	0.0	0.0
Colorado Total	9,226.2	86.9	424.6	6.5
Montana				
Broadwater	5.7	0.0	0.0	0.0
Carbon	735.0	0.0	0.0	0.0
Cascade	308.8	0.0	0.0	0.0
Fergus	210.1	0.0	0.0	0.0
Glacier	25.4	0.0	0.0	0.0
Judith Basin	90.4	0.0	0.0	0.0
Meagher	0.5	0.0	0.0	0.0
Stillwater	7.9	0.0	0.0	0.0
Montana Total	1,348.0	0.0	0.0	0.0

TABLE 14. REGION 5 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Continued)
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value Unknown</u>	<u>> 1% Sulfur</u>
New Mexico				
McKinley	113.5	0.0	0.0	0.0
Rio Arriba	3.0	0.0	0.0	0.0
Sandoval	48.2	0.0	0.0	0.0
San Juan	422.2	0.0	0.0	0.0
Valencia	0.1	0.0	0.0	0.0
New Mexico Total	1,527.2	0.0	0.0	0.0
Oregon				
Coos	1.0	0.0	0.0	0.0
Oregon Total	1.0	0.0	0.0	0.0
Utah				
Carbon	766.7	0.0	0.0	0.0
Emery	77.3	0.0	0.0	0.0
Garfield	1,011.2	0.0	0.0	0.0
Iron	5.4	0.0	0.0	0.0
Kane	1,714.5	0.0	0.0	0.0
Sevier	142.4	0.0	0.0	0.0
Unitah	40.3	0.0	0.0	0.0
Wayne	22.8	0.0	0.0	0.0
Utah Total	3,780.5	0.0	0.0	0.0
Washington				
King	44.3	0.0	0.0	0.0
Kittitas	64.4	0.0	0.0	0.0
Lewis	7.0	0.0	0.0	0.0
Pierce	65.1	0.0	0.0	0.0
Whatcom	70.4	0.0	0.0	0.0
Washington Total	251.2	0.0	0.0	0.0

TABLE 14. REGION 5 UNDERGROUND BITUMINOUS COAL RESERVE BASE BY STATE
AND COUNTY KNOWN TO FORM ACID MINE WATER ON OCCASION (Concluded)
(million short tons)

<u>State and County</u>	<u>Total Underground Bituminous Coal Reserves</u>	<u>Reserves of Listed Coal Seams (Table 9) Potentially Acid Forming</u>		
		<u>< 1% Sulfur</u>	<u>Sulfur Value Unknown</u>	<u>> 1% Sulfur</u>
Wyoming				
Carbon	39.2	0.0	0.0	0.0
Crook	1.2	0.0	0.0	0.0
Lincoln	555.9	0.0	0.0	0.0
Park	8.3	0.0	0.0	0.0
Sweetwater	3,402.8	0.0	0.0	0.0
Uinta	494.7	0.0	0.0	0.0
Weston	22.0	0.0	0.0	0.0
Wyoming Total	30,817.7	0.0	0.0	0.0

TABLE 15. NATIONWIDE UNDERGROUND BITUMINOUS COAL RESERVE BASE KNOWN TO FORM ACID MINE WATER ON OCCASION (by Region and State)

Region	Total Underground Bituminous Coal Reserves (million short tons)	Sulfur Value of Potential Acid-Forming Seams as Listed in Table 8			Percent of Total Underground Bituminous Reserves that on Occasion Have Been Known to Produce Acid Water*	
		< 1%	Unknown	> 1%	Listed Coals with Sulfur Values > 1%	Listed Coals with Sulfur Values Unknown and > 1%
Region 1						
Maryland	901.9	91.0	0.0	806.4	89	89
Ohio	17,423.3	109.4	1,747.0	13,906.8	80	90
Pennsylvania	22,788.9	968.6	2,124.5	17,928.7	79	88
Northern West Virginia	18,321.1	1,371.6	1,820.1	12,052.7	66	76
Total	59,435.2	2,540.6	5,691.6	44,694.6	75	85
Region 2						
Alabama	1,798.1	0.0	0.0	66.0	4	4
Eastern Kentucky	9,466.5	3,988.8	833.6	2,051.6	22	30
North Carolina	31.2	0.0	0.0	0.0	0	0
Tennessee	667.1	67.0	22.1	185.8	28	31
Virginia	2,833.2	211.2	10.4	381.7	13	14
Southern West Virginia	16,056.5	5,477.4	558.1	4,326.1	27	30
Total	30,852.6	9,744.4	1,424.2	7,011.2	23	27
Region 3						
Illinois	53,441.9	1,017.9	8,257.6	34,753.2	65	80
Indiana	8,948.5	157.0	679.7	4,377.8	49	57
Western Kentucky	8,719.9	0.0	207.2	7,365.8	84	87
Michigan	117.6	0.0	0.0	0.0	0	0
Total	71,227.9	1,174.9	9,144.5	46,496.8	65	78

*See footnote page 64

TABLE 15. NATIONWIDE UNDERGROUND BITUMINOUS COAL RESERVE BASE KNOWN TO
FORM ACID MINE WATER ON OCCASION (by Region and State)
(Concluded)

Region	Total Underground Bituminous Coal Reserves (million short tons)	Sulfur Value of Potential Acid-Forming Seams as Listed in Table 8			Percent of Total Underground Bituminous Reserves that on Occasion Have Been Known to Produce Acid Water*	
		< 1%	Unknown	> 1%	Listed Coals with Sulfur Values > 1%	Listed Coals with Sulfur Values Unknown and > 1%
Region 4						
Arkansas	306.0	0.0	0.0	0.0	0	0
Iowa	2,884.9	0.0	20.7	158.3	5	6
Missouri	6,073.6	0.0	596.4	1,575.4	26	36
Oklahoma	860.1	0.0	0.0	0.0	0	0
Total	10,124.6	0.0	617.1	1,733.7	17	23
Region 5						
Colorado	9,226.2	86.9	424.6	6.5	< 1	5
Montana	1,348.0	0.0	0.0	0.0	0	0
New Mexico	1,527.2	0.0	0.0	0.0	0	0
Oregon	1.0	0.0	0.0	0.0	0	0
Utah	3,780.5	0.0	0.0	0.0	0	0
Washington	251.2	0.0	0.0	0.0	0	0
Wyoming	30,817.7	0.0	0.0	0.0	0	0
Total	46,951.8	86.9	424.6	6.5	< 1	1
National Total	218,592.1	13,546.8	17,302.0	99,942.8	46	54

*These percentages represent worst case situations as can be seen by the BCR survey, Table 7, which indicates that only about 44% of the mines in Region 1, 17% of the mines in Region 2, 23% of the mines in Region 3, 12% of the mines in Region 4, and none of the mines in Region 5 produce acid mine drainage.

C. UNDERGROUND MINE DRAINAGE ABATEMENT

1. Use of Acid-formation Inhibitors: Control measures, based on eliminating one or more of the factors which influence pyrite oxidation, have been proposed to abate the formation of acid mine drainage.

a. Biological Control: As discussed in Section III A, numerous researchers have attributed increases in acid production from 2- to 17-fold to bacterial activity. The control of acid formation by applying inhibitors or biocides to retard the action of bacteria is based on reducing or eliminating this biological contribution to acid formation. Although biological control has frequently been discussed, the published experience in this area has been sparse and inconclusive. Several attempts have been made to apply inhibitors to underground mines but have met with little or no success.

As early as 1950, an experiment was conducted at West Virginia University⁶²⁵ to test the effect of acriflavine on acid formation. The Weirton mine in the Upper Freeport coal seam was selected as the test site. This mine had the advantages of an isolated location and of being a new mine where no previous acid accumulation existed. The effluent from the mine had an acidity of 400 mg/l as CaCO₃ and a pH of 4. Sulfur-oxidizing bacteria were present in the water.

In the experiment, exposed coal faces in an acid-producing section of the mine were sprayed daily for one week, then at two- to three-day intervals for two weeks with a 1:1000 solution of acriflavine, an antiseptic which was reported to prevent the growth of sulfur-oxidizing bacteria. The faces remained neutral for two weeks but, after that period, the section was again found to be producing acid. The section was cleaned and the faces sprayed again for one week. Acid production was again noted the second day following cessation of spraying. With no further explanation, the authors concluded that the bulk of acid was formed in the strata immediately overlying the coal, since the wet shale above the coal was found to be acidic. The impracticality of applying an inhibitor was demonstrated by the authors as they were unable to find even a suitable test site in a mine in the Pittsburgh seam due to the extent and complexity of the workings.

Shearer and Everson^{265,570} presented apparent evidence of the existence of bacteriophages, found in soil or compost, which dissolve (lyse) bacteria. Bacteria were inoculated with the phages under laboratory conditions and bacterial growth was inhibited for up to 7 days. Concentrated growths of bacteriophages were then isolated from neutral mine water and used to inoculate coal piles. The acid-production rate was reduced for 5 months, and further reinoculation restored inhibition. The anti-bacterial agent was later identified^{569,572} as Caulobacter. It was believed that Caulobacter may compete for the food supply and reduce the number of iron bacteria or they may attack the iron bacteria directly.

The inhibitory effect of bacteriophages was subsequently field tested at the Robena No. 2 mine by injecting 51,100 litres (13,500 gallons) of water collected from another mine into the Robena mine. The inhibitory

effectiveness of the injected water was previously demonstrated in laboratory experiments. No inhibition of acid production from the Robena mine was noted. Shearer et al. suggested that the topography prevented access of the inhibitor to some acid-producing regions. However, subsequent laboratory tests indicated that inhibitory effectiveness was absent at the time the water was collected. A second inoculation at the Robena mine was made but no change was noted in the acidity of the drainage from the test site over a period of several months. The injection of laboratory-prepared concentrates of the anti-bacterial agent into underground test mines was planned but apparently not carried out.

The work by Shearer et al. and by West Virginia University indicates the difficulty that many researchers have found in duplicating laboratory results in underground mines. Unless the exact points of acid formation are known and accessible, application of inhibitors is a fruitless endeavor. The Weirton mine used by the West Virginia University group was an ideal site--isolated and containing no previous acid accumulation. Yet the application of acriflavine to abate acid formation was not successful. Either the biocide itself was ineffective or acid may have been formed at a location remote from the test site and transported into the test area. Even under what appear to be ideal conditions, all of the parameters which affect acid formation are difficult to define.

Unresolved controversies, such as the role of sulfur-oxidizing bacteria in acid formation and the acid-forming potential of the various pyritic materials associated with coal and the surrounding strata, further complicate the problem.

The inhibitory properties of *Caulobacter*, which were used by Shearer for his studies, were questioned by Baker and Wilshire⁷⁹ who also conducted inhibition studies with the microorganisms. In a pilot-plant study, they found that *Caulobacter* had no effect on AMD production rates and were not parasitic to chemoautotrophic (those organisms able to use the energy from chemical reactions to produce nutritive substances from carbon dioxide and water) organisms.

Streptomyces aureofaciens, an antibiotic-producing bacteria, was found by Shearer et al.⁵⁶⁹ to be effective against acid-producing bacteria in test-tube cultures and on solid media.

Charnego¹⁶⁷ reported that quaternary ammonium compounds inhibited acid formation by reducing the oxygen uptake of *F. ferrooxidans*, based on laboratory tests. Charnego's results corroborated the findings of Ashmead,⁷³ who demonstrated that certain quaternary ammonium compounds in the concentration range of 5 to 6 mg/l were lethal to bacteria associated with AMD formation. The biocides were not field tested.

Walsh and Mitchell⁶⁷¹ demonstrated that ferrous ion concentrations of 100 mg/l inhibited iron-oxidizing *Metallogenium*. When *Metallogenium* was inhibited, the rate of pH decrease from iron oxidation was slowed.

In the view of the authors of this report, the use of this inhibitor, however, would increase the amount of pollutants in the water since the additional ferrous ions in mine water would create more of a problem to the coal industry than the natural situation. In addition to the ferrous iron being difficult and costly to remove from mine effluents, it can be oxidized to ferric iron, which can oxidize even more pyrite.

The reported experience with "at-source" abatement by means of biological control of acid formation has been scanty. This control technique presents several problems, and the reasons why more researchers are not pursuing this area of investigation include:

- (1) The production and use of biocides may be prohibitive as to cost of production and toxicity;
- (2) The actual points of acid formation, especially in an abandoned mine, will probably not be accessible and perhaps not even known. Therefore, the real effectiveness of any inhibitor is questionable;
- (3) Complete bacterial control would necessitate inoculating all of the infiltrating water--an impossible task;
- (4) Due to the dilution effect, periodic inoculations would have to be made to maintain or re-establish the effectiveness of the inhibitor;
- (5) Since the formation of acid mine drainage from the oxidation of pyrite is a slow, continuing process, occurring over long periods of time, any inhibitors or coatings applied for the purpose of abatement must be kept in place for equally long time periods;
- (6) Proposed methods of chemical inhibition of bacteria would result in additional pollutants which would have to be removed from the effluent;
- (7) The major problem with this control technique is that the chemical oxidation and leaching process would continue, regardless of bacterial inhibition.

In summary, the feasibility of successfully applying biological inhibitors to control acid formation is very low, and the costs would be prohibitive.

b. Coating of Pyritic Surfaces: Acid mine water is formed by the reaction of pyrite with oxidants such as oxygen and ferric ion. Whether or not water enters into the reaction is uncertain, but it is the transporting agent for the products of pyrite oxidation. "At-source" abatement of acid formation, by applying chemical coatings directly to the pyrite surface, is based on eliminating oxygen and water contact with pyrite. Very few efforts to actually utilize this abatement method are recorded in the literature.

Braley¹³⁶ reasoned that ammonia would react with iron sulfate to produce a protective coating on all oxidizable surfaces by precipitating iron as gelatinous ferric hydroxide. To test this theory, an abandoned mine was filled with ammonia gas. A slight increase, rather than the desired decrease in acidity of the mine water, was observed over the following 11 weeks.

Cotton¹⁹⁶ postulated that the oxidation of pyrite was electrolytic in nature and suggested the following: Hydrogen sulfide dissociates into sulfur, hydrogen and free electrons. The hydrogen combines with water and oxygen to form hydrogen peroxide which then combines with sulfur or pyrite to form ferrous or ferric salts or sulfuric acid. This reaction can be interrupted by electrical contact of metal salts which have a higher electrical potential than sulfur or iron. The metal salts suggested were zinc, chromium, aluminum, or phosphorus. Application difficulties were encountered, and the work was discontinued.

Phosphate, chromate, and alkali solutions and coatings were tested by Braley¹³⁷ to determine their effectiveness in preventing the reaction of sulfuritic material with oxygen. It was concluded that phosphates, chromates, and sodium hydroxide, either in solution or intimately dried on the pyritic surface, have little or no effect upon the rate of oxidation of pyrite.

Clark¹⁷⁴ also tested phosphates and chromates and reported that they did inhibit acid formation under laboratory conditions. However, he used more concentrated solutions than did Braley. Clark field-tested four chemical inhibitors: solutions of dibasic calcium phosphate, tribasic calcium phosphate, sodium pyrophosphate, and rock phosphate, which were sprayed or dusted on mine surfaces. He concluded that certain phosphates may be effective in retarding the decomposition of pyrite, but the amount of inhibiting substance needed for any particular area had not been determined.

Bloom *et al.*¹¹¹ reported on mono-, di- and tri-chloro derivatives of silane (SiH_4) for treatment of sulfide materials. The compounds reacted in the gaseous state with sulfides to form a hydrophobic surface. Bonding was stable in acid media. However, the gaseous substances also reacted with water. Field testing was not completed.

Tyco Laboratories^{577,655} applied a silica- and aluminum/silica-gel treatment process to refuse piles. The silica gel endured for 305 centimeters (120 inches) of equivalent rainfall and the alumina/silica for 1,270 centimeters (500 inches). The gel was not tested in underground mines.

Peeler⁵¹⁹ discussed treatment of soil containing iron pyrites with an aqueous alkali metal silicate solution. The silicate solution reacted with sulfuric acid to form a silica gel.

Flynn²⁷⁸ described a method of inhibiting the rate of pyrite oxidation by applying a mixture of pulverized phosphate and sulfuric acid. The phosphate rock reportedly converted any soluble iron compounds to non-acid forming phosphates which coated the unreacted iron sulfides, thereby eliminating air and water contact.

"At-source" abatement by coating acid-forming materials with an impervious shield to eliminate air and water contact with pyrite surfaces is not practical due to the inaccessibility of the enormous volumes of fractured coal and rock that contribute to acid formation. New pyritic surfaces are constantly being exposed as a result of roof falls and pillar collapse. This means periodic applications would have to be made to effect complete control. This would represent an impossible problem due to the inaccessibility of mined-out sections. The additional pollutants that may be formed from any inhibitor material would also compound the problem.

2. Hydrologic Controls

a. Water-infiltration Control: Water may enter the underground mine through surface fractures, permeable overburden, subsidence areas, bottom clay, adjacent non-regraded surface mines, and shaft, drift, slope, and borehole openings. To permit active mining, this water normally must be discharged from the mine via pumping or by gravity drainage. Since any available products of pyrite oxidation may also be transported from the mine, the mine water may require treatment before being discharged to the receiving stream.

Water-infiltration control techniques are designed to reduce the quantity of water that enters an underground mine, thereby reducing the volume of water that must be treated and discharged. If the volume of influent water can be substantially reduced, some savings in pumping and treatment costs would accrue to the coal industry.

Methods proposed, but not demonstrated,⁴⁶⁰ for determining sources, paths, and quantities of water entering underground coal mines include:

- (1) Photogeologic methods--most applicable to mining is small-scale and medium-scale aerial photography to map geological structures such as folding and fracturing of rock strata which are fundamental to (a) the occurrence of water in an underground mine, (b) the routes the water follows once it enters the subsurface, and (c) where the water will discharge.
- (2) Field examination--to verify all structures previously inferred during the photogeologic studies. If conducted by an experienced hydrogeologist or engineering geologist, field examination can suggest many possibilities for avenues of water infiltration into an underground mine.
- (3) Geophysical methods--the most applicable to coal mining are borehole techniques. Use of borehole methods facilitates quantitative measurement of material properties (density, porosity, permeability, and elasticity) and provides qualitative information such as zones of water influx, relative clay content, and lithology. Other less-applicable geophysical

methods include surface, airborne, and in-mine techniques. Geophysical methods can be used to locate geologic and man-made situations--joints, faults, sand intrusion into a coal seam, unconfined aquifers, abandoned mine openings and workings--all of which contribute to the problem of locating and quantifying potential sources of water entering underground coal mines.

(4) Subsurface geologic methods--measurements and observations of rock lithology; stratigraphy; gross structural geological characteristics of the overburden; and extent, thickness, and rank of coal are made by means of diamond core drilling, drilling and geologic logs, and laboratory testing. Individual geologic maps can be overlaid and used as a predictive tool to define slight to severe water inflow problem areas.

(5) Hydrogeologic methods--information about hydrogeological characteristics of rock units can be made by observations and measurements in existing wells, boreholes associated with coal exploration and development, and adjacent mines. Specific techniques include seepage tests, pressure tests, pump tests, and water analysis. The data obtained pertain to (a) the presence and extent of aquifers and aquicludes, (b) groundwater levels and pressures, (c) hydraulic parameters, and (d) water quality.

(6) Tracer methods--artificially-injected and naturally-occurring tracers such as organic dyes, inorganic substances, and radioactive isotopes have been used to physically trace movements of water which may enter an underground mine. No single tracer is ideal for all situations, and the applicability of this method is limited to small areas.

Control of surface-water infiltration into abandoned deep mines has been implemented in several "at-source" abatement programs under Operation Scarlift in Pennsylvania.

The Catawissa Creek Project⁷⁴⁵ involved the reconstruction of a streambed to prevent the stream from entering the underground mine. With an average of 21.88 million litres (5.8 million gallons) of water per day prevented from passing through the mine, the load of acid to Catawissa Creek was reduced by an average of 830 kilograms (1,830 pounds) per day. However, the amount of iron discharged into the creek was approximately the same because the reduced volume of flow carried a higher concentration of iron.

Water-infiltration control was accomplished in the Beech Creek Stream Project⁷⁴⁵ where a significant portion of the streamflow of Little Sandy Run was lost to deep mines through subsidence areas. A new channel was constructed and the subsidence areas filled to achieve positive drainage into the channel. An estimated 1.5 million litres (0.4 million gallons) per day of water was prevented from entering the mines. Thus, an average of

283 kilograms (624 pounds) of acid and 61 kilograms (156 pounds) of iron per day, which would have been carried by this amount of flow, were not discharged into the stream.

In the Tioga River area,⁷⁴⁵ reconstruction of a streambed and restoration of strip mines prevented an estimated daily infiltration of 4.66 million litres (1.23 million gallons) per day of water into underlying mines. This reduction in flow was estimated to result in an average daily reduction in acid load to the stream of 3,850 kilograms (8,480 pounds).

Surface sealing of subsidence areas and water diversion were used in conjunction with deep-mine sealing at the Shaw Mines Complex to realize a 45-percent reduction in acidity. At Alder Run, a channel was relocated and diverted water away from deep-mine openings.²⁸³

Techniques to seal surface cracks, rechannel surface water, and seal boreholes and mine openings have been successfully demonstrated at numerous other sites than those listed above. These sites include many active mines.

Water-infiltration control might also be achieved by intercepting ground water before it enters the mine. This method of acid mine drainage abatement has been discussed extensively but is still in the theoretical stage. One such plan¹⁵ was proposed to eliminate the acid discharge from the Todd mine into the Lake Hope drainage basin. An investigation of the area suggested that there were three main aquifers: a shallow water table above the coal; below the coal, a brackish aquifer between 300 and 400 feet; and a deeper fresh-water aquifer. In the proposed plan, the overlying aquifer would be connected to either of the lower aquifers by weeping wells so that water would bypass the coal and related strata instead of percolating through them to pick up pyrite oxidation products.

Loofbourow and Brittain,⁴²² in discussing the conditions favorable to dewatering, noted that: aquifers should be predictable as to position and productivity; ground should have some minimum coherence; comparatively shallow aquifers are developed more rapidly and are developed and pumped at less cost; ground should be easily bored; and water flows should not be extreme.

Considerable hydrogeological data is required to determine the feasibility of this approach to infiltration control. According to Parizek⁷⁵³ these data include the spatial distribution, thickness, and geometry of aquifer and non-aquifer units; hydraulic boundaries that either restrict the flow of ground water (stratigraphic pinchouts, fault offsets, erosional unconformities) or serve as recharge sources (channel sandstones, flooded deep mines, fractured roof rock etc); permeability and storage distribution; infiltration capacity of streambed sediments; vertical permeability of confining beds; and hydraulic heads among and between various aquifer and non-aquifer units.

In some instances, and it is the opinion of the coal industry that such opportunities are limited, subsurface-infiltration control may reduce the

volume of water entering the mine. Such instances could include all types of underground mines during development and deeper, below-drainage mines during the entire life of the mine. Normally, ground water that enters underground mines above drainage has not migrated through aquifers for any appreciable distance but has originated as precipitation or surface-water flow over the ground surface directly overlying the mine. As mining progresses, secondary fracturing of the overburden will result, which will enhance the flow of this water into the mine. It is difficult to believe that under normal circumstances it will be technically or economically possible to drill a sufficient number of wells to achieve any significant infiltration reduction. A pilot dewatering project to demonstrate the technical and economic feasibility of dewatering an active underground mine to control or prevent acid mine drainage is currently being conducted in central Pennsylvania by W. A. Wahler and Associates. The results of this demonstration project should be critically evaluated, particularly as they relate to developing mines and mines below drainage, to determine whether additional research on dewatering techniques is advisable.

b. Water Handling: The most effective abatement procedures that can be utilized in an operating mine involve the handling of water once it gains entrance to the mine. The principles involved, as described in Principles and Guide to Practices in the Control of Acid Mine Drainage (Attachment A), are minimizing the contact time between the water and acid-forming materials and removing the water as expeditiously as possible. Minimization of the contact time also assumes that care will be exercised in the placement of acid-forming materials.

- (1) All gravity drains and sumps should be maintained in such a manner that they are kept free of acid-forming materials.
- (2) If gravity drains are constructed in material that is acid producing, consideration should be given to enclosing the flow.
- (3) Highly pyritic coal-cleaning refuse normally should not be returned underground but should be disposed of on the surface where positive action can be taken to eliminate any acid discharge. Underground stowage should only be considered at those locations where the mine is essentially dry or at those mines or sections of mines where such wastes will be completely inundated.
- (4) Pumping schedules should be arranged to prevent large deviations in the water level within the mine. If acid-forming materials are exposed, the cyclic wetting and drying will result in an acceleration of the acid-formation process.
- (5) Major sump locations should be determined prior to mining and consideration of procedures for accomplishing expeditious dewatering should be an integral part of the pre-mining planning.
- (6) As mining progresses, secondary sumps should be constructed and positive pumping established to prevent uncontrolled water movement through the mine.

These procedures are fairly well known and have been put into practice by most major coal-producing companies. Comprehensive studies to evaluate these procedures have not been conducted, and it is difficult to accurately assess their effectiveness. However, it can be inferred, from the data from the BCR and EPA surveys, that the alkaline quality of many discharges originating from mines in acid-producing seams was due to implementation of these procedures. Coal-industry experts caution that, although acid production can often be significantly minimized by implementing these practices, it cannot always be eliminated.

Additional reductions in acid mine drainage pollution can be achieved through wider use of these practices. To identify and promote the use of such practices, it is recommended that the Bureau of Mines prepare a manual on control of acid mine drainage and develop a program that will result in the dissemination of this information throughout the coal industry. It would seem appropriate to utilize the Bureau of Mines State Liaison Officers in the dissemination effort. It would also seem appropriate to provide them with complete information concerning the mine-drainage problems in their respective states. By doing this, the coal industry in any given state would have a single source of complete information that they could utilize.

3. Mining Methods: The most common past practice has been to mine coal from the outcrop rather than driving slope or shaft entries. Such mines are usually driven up-dip wherever possible to, in part, facilitate gravity drainage and reduce pumping costs. When abandoned, it is extremely difficult to seal these mines and eliminate the discharges because of the large hydrostatic heads that will develop as the mine floods. Down-dip mining, which would eliminate all gravity discharges, therefore, has been suggested^{512,536} as a viable method of guaranteeing the elimination of the mine discharge when the mine is abandoned and sealed.

An evaluation of up-dip versus down-dip mining was conducted for EPA by Skelly and Loy Engineers and Consultants.⁴⁵⁷ Water quality was evaluated at a pair of abandoned mines--one developed up-dip and one developed down-dip. Since the mines were apparently similar in all other respects, the primary factor controlling water quality was the direction of mine development. It was determined that the acidity concentrations in the main discharge from the abandoned mine developed to the rise were consistently in excess of 2200 mg/l, while those of the discharge from the mine developed to the dip were generally less than 100 mg/l.

An active mine with units operating up-dip and down-dip was evaluated, as part of the same study, to ascertain major advantages and disadvantages of each method. The authors concluded: down-dip mining at the active site was no more or less productive than up-dip mining; pumping costs may be substantially increased by down-dip mining but they will probably not reach the point of adversely affecting production economics; despite extremely low conveyor-belt-haulage unit cost, percentage cost increases, incurred by haulage up-grade rather than down-grade, range from a few to several hundred percent; and effective sealing of mines in steeply dipping seams can only be assumed where mines have been developed to the dip and the lower outcrop or barrier integrity has been maintained.

In view of the prospect that mine operators might have to treat the mine discharge for many years after active mining has been completed, the additional costs incurred by mining down-dip may be offset by elimination of treatment costs through effective sealing upon abandonment. In acid-producing seams, down-dip rather than up-dip mining might be considered whenever the option exists of using either method. One technological limitation to the successful application of this method is the problem of estimating the required thicknesses of barrier pillar and coal outcrop so that coal recovery is maximized while outcrop or barrier failure is minimized.

Longwall mining has been suggested as a mining method that possibly will result in the inhibition of acid mine drainage formation.⁵⁶⁶ It has been theorized that this acid reduction might result from several circumstances. First, if the acid condition normally experienced when mining a particular coal bed is due to pyrite found in the coal, the availability of such pyrite will be greatly reduced, since nearly 85 percent coal recovery is attained by longwall mining. The second circumstance concerns the effect of caving which would essentially be complete and uniform. It is theorized that caving would: (a) reduce the volume of water which would come in contact with oxidized materials; (b) restrict water flow through the mined-out area; and (c) limit the amount of air that would come in contact with any pyrite that is exposed or is available for oxidation.

The effect of longwall mining on water quality should be exhaustively studied as this method of mining has the appealing advantages of increasing production and of recovering a larger percentage of our valuable coal resource. The results of such a study might indicate that total roof collapse is the circumstance that results in improved water quality. If so, achieving roof collapse could be an abatement measure that would be recommended as an objective when using other mining systems.

4. Mine Closure

a. Air Sealing: The use of mine seals to exclude air from abandoned mines, thereby decreasing oxidation and the formation of acid mine drainage, has been the subject of many investigations since the early 1900's.

The initial research on sealing mines to reduce acidity was conducted in the 1920's and early 1930's by the Bureau of Mines.^{413, 415, 416, 418} It was observed that the sealed mines were not as acid as adjacent mines which had not been sealed. As a result of these studies, a program to seal abandoned mines in ten states was initiated in 1933 under the Civil Works Administration (CWA). Pennsylvania, West Virginia, Alabama, Ohio, Kentucky, Maryland, Tennessee, Virginia, Illinois, and Indiana were involved in the program.

Three types of mine seals were employed in the sealing operations.³³⁴ Dry seals, constructed of earth, brick, concrete, or stone, were used to exclude air from openings from which no drainage flowed. Openings from which drainage flowed were closed with air seals. This type of seal excluded air but permitted the normal outflow of water. In some cases, the only effective

method of sealing was to flood the workings by building a concrete barrier to the full height of the drainage openings. The hydrostatic pressure developed by such a construction was normally relieved by removing the mine water at a point above the roof of the mine.

The CWA program was dissolved in March, 1934, but the mine sealing was continued as state projects in Ohio, West Virginia, and Pennsylvania.^{518, 641, 642}

In 1935, a large-scale mine-sealing program was inaugurated under the direction of the Works Progress Administration (WPA).⁴⁷⁷ The objective was to reduce the acid flowing into streams of Pennsylvania, Ohio, Kentucky, and West Virginia. This undertaking was a continuation of the work intitated by the CWA.

Several investigators^{165, 329} have reported on the mine-sealing projects conducted during the 1930's. These reports indicate that the seals resulted in a reduction of acidity. Fellows,²⁷⁶ for instance, noted reductions of 50 to well over 90 percent in the "sulfuric acid" flow from abandoned mines. However, the effectiveness of the mine-sealing programs was not long lasting as many of the seals either failed, due to lack of maintenance, or were broken in order to resume mining during the war years in the 1940's. Several more recent studies^{9, 491, 722} were undertaken to assess the effectiveness of these projects but all concluded that such an assessment was impossible due to the lack of information.

It should be noted that most of the early mine-sealing projects were depression years "make work" projects. Although many highly qualified engineers and skilled artisans were involved, the work performed, using the limited technology and materials available, was not completely satisfactory in view of the demanding and thorough job that is required to air seal a mine. Any improvement in water quality, and there are many examples, probably was the result of mine flooding and inundation of acid-forming materials, and of reduction in discharge flow rates, rather than oxygen reduction, which was the objective of air sealing.

In a more recent study, Braley¹²⁶ evaluated the effectiveness of air sealing seven abandoned drift mines. In all cases, the mines had gravity drainage and the overburden thickness ranged from 15 to 61 meters (50 to 200 feet). Criteria for determining the effectiveness of air sealing were: (1) reduction in the amount of acid produced, and (2) decrease of oxygen concentration in the mine atmosphere. He concluded that sealing of abandoned drift mines, when the coal seam was above drainage level, was ineffective in reducing the quantity of acid which was discharged. There also was no evidence that sealing of the mine substantially reduced the oxygen content of the atmosphere inside the mine.

In order to evaluate the effectiveness of air sealing, the Bureau of Mines⁴⁸³ sealed a 31-hectare (77-acre) above-drainage coal mine. One dry seal and seven wet seals were constructed. Mine-atmosphere samples were collected

from five locations after sealing. A reduction in acidity of 150 mg/l shortly after sealing was directly attributed to the effects of sealing. The mean total acidity was 514 mg/l before sealing and 211 mg/l for almost 3 years after sealing. The oxygen level decreased from 20.9 to 17.0 percent following sealing. Five months later, the oxygen concentration began to fluctuate and did not decrease below 16 percent.

Better overall results would have been obtained had it been practical to find all permeable zones and, if suitable proven techniques were available, to seal them.

Several small underground mines, which were discharging acid mine drainage into Taylor Run, Red Run, and Fishing Hawk (all tributaries to Shavers Fork), were air sealed by the West Virginia Department of Mines. Evaluation of the effectiveness of the air sealing, which was conducted by EPA,⁵⁶⁴ was limited because either insufficient or unreliable data were collected prior to sealing. However, the overall results indicated that the concentrations and load discharging into Shavers Fork did not change to a great extent in the two years following sealing.

Active research on acid mine drainage has been conducted at the McDaniels Test Mine since 1964. This pilot-scale facility is intermediate between laboratory scale and demonstration scale. The effect of oxygen concentration on acid production was studied⁵²⁸ by maintaining the oxygen concentration of the McDaniels Test Mine at less than 1 percent for one year. Acid loads decreased to about 60 percent of the values for the previous year. This was approximately one-half of the expected decrease if steady-state conditions were reached.

It was estimated that if all oxygen could be excluded from the mine, 4-1/2 years would be required to remove 90 percent of the oxidation products present at the time oxygen was first eliminated from the mine. This indicates the care that must be exercised in interpreting mine-drainage field data because of the slow response of natural systems to imposed environmental changes and the erratic output due to hydrologic variations. Conclusions cannot be made on a single mine's discharge data over a 1- or 2-year period.

Both surface reclamation and underground mine sealing were carried out during the Elkins project.³⁵⁶ Eleven "wet" seals (designed to allow water to leave the mine but prevent air from entering) were constructed in an 809-hectare (2000-acre) underground mine complex and one seal in a small, isolated mine. The sealing of the large mine was not completed and subsidence over large portions of the mine was not corrected. All known openings to the small, isolated mine were sealed.

The oxygen level inside the large mine was the same as the air outside the mine. The acid-load data showed little if any improvement.

The oxygen content of the small, isolated mine decreased to 11 percent during the first two years following sealing, and then increased to and remained at about 15 percent. The acidity, iron, and sulfate concentrations decreased but an increase in flow occurred so pollution loads showed little change.

Conclusions of the project were that air sealing of underground mines to eliminate oxygen could not be accomplished under the conditions encountered at Elkins. Even under the best conditions, oxygen was reduced to only 7 percent.

R. D. Hill,³⁵⁵ when reviewing several mine-sealing projects, emphasized that in two projects, where extraordinary efforts to seal all known openings, subsidence, etc., were used, there was still an average of 15 to 17 percent oxygen in the mine atmosphere, indicating that air still had access to the mines. While the reduction of acidity was about 50 percent, part of that reduction could be attributed to preventing water from entering the mines. Hill's conclusions were: that 50 percent effectiveness was all that could be expected even from the best air-sealed mine and even then maintenance would be necessary to assure the integrity of the seal; and that mines with shallow cover, where outcrops were surface mined and which were highly subsided, would be even less conducive to air sealing.

In a survey conducted by Bucek *et al.*⁷²² pre- and post-closure water quality and quantity data were collected for 84 underground coal mines. They concluded that air sealing reduced acidity concentrations on the average by 50 percent. It was also pointed out in this study that air access was not eliminated even under the extraordinary measures that were taken during the sealing projects in Pennsylvania and West Virginia. All of these mines had shallow overburden (22-24 meters or 72-79 feet). A mine, having an overburden averaging 144 meters (472 feet), that was air sealed in West Virginia in 1972 was also evaluated. At this site, the pollutant concentrations increased considerably after sealing but have been diminishing at considerable rates since then. Based on this finding, they suggested that mines with thicker overburden should be more suitable for effective air sealing.

The major problem with air sealing underground mines is the elimination of air access through the fractured and porous overburden. With every change in atmospheric pressure, the mine "breathes," thereby replenishing the oxygen in the mine. As was previously noted in reviewing the mine-sealing work done by EPA at Elkins, West Virginia and by the Bureau of Mines, even when extreme precautions were taken to eliminate air access, the oxygen content of the mine did not drop below 7 percent. From research conducted at the McDaniels Test Mine on pyrite oxidation, it was estimated that the oxygen concentration would have to be less than 1 percent to effect a greater than 90 percent reduction in acidity.

The demonstrated effectiveness of air sealing in reducing acid loads is only about 50 percent. To ensure even this limited effectiveness, periodic maintenance of the seals is required. Additionally, subsidence and fracturing

may open new air access routes which would mean additional surface sealing. Considering that air sealing has been demonstrated to be only moderately effective in reducing acidity, and that there are major limitations inherent to its successful application, further research in this area appears impractical.

The feasibility of blanketing an abandoned deep mine with an inert gas in order to eliminate acid mine drainage formation was evaluated by Cyrus Wm. Rice Division, NUS Corporation.⁵⁵² The results of the study indicated that, although positive differential pressures could be maintained within the two mines investigated, continuous gas injection was necessary.

Mines with shallow overburden or with porous or fractured confining strata would not be suited to inert-gas injection since the gas would escape and, with each change in barometric pressure, oxygen could enter the mine. Periodic maintenance of the seals would be necessary, and fracturing and subsidence would open up new pathways through which gas could escape. This abatement method is not economically feasible since it requires continuous operation and, in this respect, is not a permanent solution to the acid mine drainage problem.

b. Hydraulic Sealing: Laboratory investigations and field studies have demonstrated that oxidation of pyrite can be eliminated by covering pyritic materials with water.

Braley¹²⁸ reported a significant improvement in water quality as a result of deep-mine flooding. He stipulated that the extent of beneficiation is dependent upon extent of total flooding as determined by coal and surface elevations.

The coal industry recognized the importance of mine flooding and, through the Coal Industry Advisory Committee (CIAC) to the Ohio River Valley Water Sanitation Commission (ORSANCO), published in 1964 the manual Principles and Guide to Practices in the Control of Acid Mine-Drainage.⁵³⁴ One of these control measures called for effective planning of mine closing, including sealing, to minimize the formation and discharge of acid mine drainage.

Until the 1960's, most of the seals were constructed of cinder block and cement, which deteriorated in acid waters and, therefore, proved ineffective. As a result of the increased interest in environmental control, new mine-sealing techniques had to be developed. Today, hydraulic deep-mine seals are constructed of concrete and grout⁶³⁹ and are designed to withstand water pressure, prevent inflow, and contain water.

From 1966 to 1969, the Halliburton Company^{270-273, 499} conducted a study to develop and field test new methods for watertight seals and bulkhead construction. Based on the results of field tests, they concluded that a mine seal, composed of front and rear bulkheads of a quick-setting slurry with expansive-type cement filler between, would be satisfactory to seal a mine drift to hold the resultant head of impounded water.

In the last ten years, source-abatement methods have been employed in a number of government-sponsored programs. In the Moraine State Park²⁸⁴ watershed area, 22 abandoned deep mines were sealed by installing 69 hydraulic mine seals of double-bulkhead construction, grouting, and surface sealing. Prior to the abatement, over 85 percent of the acidity, draining to what is now Lake Arthur, came from these 22 mines. The deep-mine sealing and grouting work resulted in 68-percent reduction in net acidity. Water quality has remained constant since construction was completed in 1971.

Mine-drainage abatement projects²⁸³ also have been completed in the Argentine and Whiskerville mines, in the Shaw Mines Complex, and in the Ernest Complex in Pennsylvania. The Argentine and Whiskerville project consisted of installing 32 hydraulic deep-mine seals. Six deep-mine observation holes were installed to monitor the effect of sealing on water quality. As a result of the sealing operation, the flow was reduced by 70 percent and the net acidity decreased by 85 percent. Abatement work in the Shaw Mines Complex included installing five deep-mine hydraulic seals and grout curtains. Although the effect of deep-mine sealing could not be directly evaluated, the completed work resulted in a 45-percent reduction in acidity. Mine sealing completed in the Ernest Complex resulted in a 78-percent reduction in acidity. Based on these three projects, it was concluded that deep-mine sealing projects were the most effective of all abatement methods in reducing acidity.

The environmental effectiveness of close-down procedures at 84 sites throughout the eastern United States was evaluated by Bucek *et al.*⁷²² They reported that double-bulkhead sealing resulted in 45- to 99-percent reductions in acidity at 80 percent of the sites visited.

Klingensmith⁷⁴⁵ reported on the results of using clay seals to achieve inundation of an abandoned deep mine. Prior to sealing, the discharge amounted to 354,276 litres (93,600 gallons) per day and contributed 246 kilograms (543 pounds) per day of acid to the stream. Several months after construction was completed, only a small trickle was found in the area of the former discharge.

Mine flooding has been demonstrated to be the only viable method of minimizing or eliminating acid discharges from underground mines following abandonment. Since such flooding will occur naturally in those mines that are situated below drainage, the concern for developing abatement procedures must be concentrated on those mines located above drainage. Flooding will also occur naturally in drift mines that are worked down-dip if there are no unsealed openings to the surface at elevations that are lower than that of the portal. Flooding of those mines having gravity discharges is the most difficult to achieve although some success has been reported.

Successful flooding of any above-drainage mine depends on: (1) having and maintaining a stable and impermeable outcrop and/or property barrier; (2) constructing durable and impermeable seals; (3) maintaining the long-time integrity of the seals; and (4) locating and sealing all openings to the surface that have resulted from mining. For instance, it can be theorized that, if there is sufficient hydrostatic head, in shallow-cover mines, discharges could be anticipated through the overburden cover at those points where major subsidence has occurred.

It is apparent that mine closure must be considered when planning and developing a mine. The width of the barriers to be left in place must be determined, portals must be planned and developed in such a way that successful sealing can be achieved, down-dip mining should be considered if the option exists, interior section- or panel-sealing should be considered, openings to the surface should be limited, and programs to record subsidence areas should be initiated.

Flooding appears to be the best method of abating acid mine drainage formation and discharge. However, before mine flooding can be unequivocally recommended, several problems must be addressed and additional research must be conducted.

First, tremendous volumes of water will be contained and, inadvertently, by man's activities, such as highway construction, or through natural deterioration of seals and outcrop, may be released with catastrophic results. If mine-flooding becomes widespread, the Bureau of Mines should accept the responsibility of mapping these mines and monitoring all surface activities that might interfere with the integrity of the system.

Second, the quality of overburden ground water may be affected at those sites where a large hydrostatic head results from mining. Future Bureau research should determine whether or not ground-water pollution might result from sealing.

Third, a considerable quantity of coal will be wasted when leaving outcrop and barrier pillars. At present, there are no valid specifications for determining the minimum pillar size required. There is a legitimate need for the Bureau of Mines to conduct research in this area. The pillar size requirements to provide adequate support for surface water bodies should also be investigated. Such research is needed since currently proposed requirements for support pillars are so stringent that longwall mining will be precluded in many areas in order that complete support be provided for even those streams that are classified as only wet weather tributaries.

Fourth, most mine-sealing programs have been conducted at abandoned mines and the conclusions drawn have been based on incomplete information concerning the mines and premining and presealing conditions. Restrictions in the location, type, and construction of the seals have been imposed by the conditions found at the site. There is also no assurance that all openings have been located and that all barriers are competent. The Bureau of Mines should undertake a mine-sealing research program in conjunction with the closure of a limited number of operating mines to determine the true effectiveness of sealing. There are, however, several recently completed, abandoned-mine-sealing programs for which adequate background information is available. There appears to be a lack of funds to continue monitoring these sites and, as much information can be gained from these experiences, the Bureau of Mines should sponsor a project to provide the required monitoring, maintenance, and evaluation.

Fifth, in certain states, section sealing of abandoned portions of active mines may be a problem due to conflicting regulations. For instance, Section 247 of the Mining Laws of Pennsylvania requires that "in a gassy mine all workings which are abandoned after the effective date of this act, or the dates such mine became a gassy mine, whichever is later, shall be ventilated." Section 291 states that "the superintendent shall not permit the mining of coal within fifty feet of an abandoned mine or any abandoned portion of any mine containing a dangerous accumulation of water, until said danger has been removed by driving a passageway to tap and drain off said water." Performance of either of these safety requirements would prevent section sealing or would eliminate its effectiveness. That section sealing is effective has been demonstrated at least at one site. At the Saginaw mine²⁶⁶, as a result of sealing abandoned sections, only minimal amounts of water were found within the active sections of the mine. More extensive investigations into the effectiveness of section sealing should be conducted, and the conflict between existing mining laws and the requirements for section sealing should be resolved.

IV. RESEARCH NEEDS AND PRIORITIES

A list of potential research projects, or lines of inquiries, that were suggested by this review can be found in Table 16. Each project has received a value judgment, using a scale of 0 to 10, with 10 being the highest mark given in each of four categories. These categories include: (1) Value of This Research Being Pursued--These ratings reflect the perceived overall value of the project without relating it directly to underground acid mine drainage abatement, or restricting future research to that to be conducted by the Bureau; (2) Probability of Leading to Development of Abatement Procedures--These ratings reflect the probability that the recommended project will result in the development of new mine-drainage procedures or will better define those procedures already available; (3) Value to the Coal Industry--These ratings are based on discussions with coal industry experts and reflect what they perceive to be their needs; and (4) Value of Bureau of Mines Pursuing This Research--These ratings are based on the ratings shown for the first three categories and, in addition, reflect both the need and the benefits to be derived from conducting the recommended project.

It is recommended that, to develop "at-source" abatement techniques, the Bureau of Mines actively pursue potential research projects (6) Development of Mine Closure Technology, (10) Development of Guidelines for Implementation of Surface Water Infiltration Control Procedures, (14) Study of the Effect of Roof Collapse on Mine Water Characteristics, (16) Evaluation of the Long-term Effects of Flooding Underground Mines Above Drainage, and (17) Investigate the Effectiveness of Section Sealing in Active Mines.

A. PROJECT 6, DEVELOPMENT OF MINE CLOSURE TECHNOLOGY

Objective:

To provide the mining industry with the technology required to effectively seal a mine to ensure that all mine-water discharges are eliminated, or that flooding of a significant portion of the abandoned mine is achieved. Mine sealing must be considered an integral part of the active mining operation as the effectiveness of the sealing program will be dependent on the precautions taken during mining. Project 16 should be completed before work is initiated on this project.

Work Tasks:

(1) Determine the specifications for minimal size of all coal barriers, both internal and outcrop, to be left in place. These specifications should be dependent on projected hydrostatic head, structure of coal, competency of overburden, relationship of mine to adjacent mines in the same or other coal seams, and should ensure maximum coal recovery;

(2) Define the materials and procedures required for grouting less than minimal size barriers and fractured overburden to ensure their integrity; and

TABLE 16. LIST OF PROJECTS THAT MIGHT BE CONSIDERED FOR FUTURE RESEARCH

Potential Research Projects	Value Judgment 0-10 (10 is highest)				Comments
	Value of this Research Being Pursued	Probability of Leading to Development of Abatement Procedures	Value to Industry	Value of BOM Pursuing this Research	
1. Studies to Determine Exact Mechanisms by Which Chemical Oxidation of Pyrite Occurs	1	1	0	0	Information gaps exist that might be solved by further research. However, probability is remote that such research would lead to development of abatement procedures.
2. Studies to Precisely Define the Role of Bacteria in the Formation of Acid Mine Water	2	1	0	0	Information gaps exist that might be solved by further research. However, probability is remote that such research would lead to development of abatement procedures.
3. Develop a Method for Determining the Acid-forming Potential of Coal and Adjacent Stratigraphic Measures	3	0	3	3	Such a method might be valuable when preplanning a mine to determine the quality of water that will be generated, but research in this area will not lead to development of mine drainage abatement procedures.

TABLE 16. LIST OF PROJECTS THAT MIGHT BE CONSIDERED FOR FUTURE RESEARCH (Continued)

Potential Research Projects	Value Judgment 0-10 (10 is highest)				Comments
	Value of this Research Being Pursued	Probability of Leading to Development of Abatement Procedures	Value to Industry	Value of BOM Pursuing this Research	
4. Development of a Laboratory Procedure, that Simulates the Conditions Which Exist in an Underground Mine, for Producing Acid Water	3	1	0	1	Reliable procedures for acid generation in the laboratory are not available. Such procedures should be developed before any laboratory studies involving mine acid formation are initiated.
5. Develop a Simple Model for Predicting Mine Water Quality and Quantity	5	1	5	5	Such a model would be of value to the coal industry and would indicate the benefits to be derived from the implementation of "at-source" abatement procedures at a given site.
6. Development of Mine-closure Technology	10	10	10	10	This assumes that the objective is partial or complete flooding of mines above drainage. Prior to conducting research in this area, thoughtful consideration should be given to the potential hazards of such flooding.

TABLE 16. LIST OF PROJECTS THAT MIGHT BE CONSIDERED FOR FUTURE RESEARCH (Continued)

Potential Research Projects	Value Judgment 0-10 (10 is highest)				Comments
	Value of this Research Being Pursued	Probability of Leading to Development of Abatement Procedures	Value to Industry	Value of BOM Pursuing this Research	
7. Studies to Determine Effect of Rock Dusting on Mine Water Quality	2	2	3	3	It is speculatively concluded that rock dusting will have a positive effect on mine water quality. This effect should be considered when evaluating abatement procedures.
8. Evaluation of Materials to Inhibit Biological Oxidation of Pyrite	0	0	0	0	Applying and maintaining the effectiveness of these materials appear to be impossible tasks, and additional research is not recommended.
9. Development of Materials to Inhibit Chemical Oxidation of Pyrite	2	1	3	2	Results of this research would be applicable to surface disposal of coal-cleaning refuse. Indirectly, this could be considered "at-source" abatement as this refuse would not be returned underground.

TABLE 16. LIST OF PROJECTS THAT MIGHT BE CONSIDERED FOR FUTURE RESEARCH (Continued)

Potential Research Projects	Value Judgment 0-10 (10 is highest)				Comments
	Value of this Research Being Pursued	Probability of Leading to Development of Abatement Procedures	Value to Industry	Value of BOM Pursuing this Research	
10. Development of Guidelines for Implementation of Surface Water Infiltration Control Procedures	5	5	7	5	Many such methods have been demonstrated, primarily at abandoned mine sites, but need to be presented in manual form that can be used by industry. New specifications are needed for support pillars that will permit greater coal recovery.
11. Development of Methods for Reducing Ground Water Entry into Underground Mines	1	1	1	0	Research has been initiated by EPA. Follow-up research should be conducted by same agency to avoid duplication of effort.
12. Prepare a Manual on Mine Drainage Abatement Procedures	5	2	7	5	Such a manual would benefit the many small operators. The value of the benefit would depend on the mechanism established by Bureau to disseminate information.

TABLE 16. LIST OF PROJECTS THAT MIGHT BE CONSIDERED FOR FUTURE RESEARCH (Continued)

Potential Research Projects	Value Judgment 0-10 (10 is highest)				Comments
	Value of this Research Being Pursued	Probability of Leading to Development of Abatement Procedures	Value to Industry	Value of BOM Pursuing this Research	
13. Develop a Program to Disseminate Information on Mine Drainage Control to Coal Industry	5	2	7	5	
14. Study of the Effect of Roof Collapse on Mine Water Characteristics	7	7	7	7	
15. Develop Methods of Air Sealing to Reduce Oxygen Content in Mine	1	1	1	0	Effective air sealing is not compatible with shallow-cover mines where maximum coal recovery has been achieved. Deeper mines, where oxygen cannot enter the mine via subsided areas or secondary overburden fractures, are normally below drainage and will naturally become flooded when the mine is abandoned.

TABLE 16. LIST OF PROJECTS THAT MIGHT BE CONSIDERED FOR FUTURE RESEARCH (Concluded)

Potential Research Projects	Value Judgment 0-10 (10 is highest)				Comments
	Value of this Research Being Pursued	Probability of Leading to Development of Abatement Procedures	Value to Industry	Value of BOM Pursuing this Research	
16. Evaluation of the Long-term Effects of Flooding Underground Mines Above Drainage	10	10	10	10	
17. Investigate the Effectiveness of Section Sealing in Active Mines	7	3	7	7	

(3) Develop design specifications for constructing a mine seal that will withstand the anticipated hydrostatic head; that can be effectively linked to the roof, rib, and bottom; and that will have maximum durability.

Applicability:

Results will be applicable to future underground mines to be opened in coal seams that represent approximately 25 percent of the reserves in the Northern Appalachian Region and 10 percent of the reserves in the Southern Appalachian Region.

Sequential Restraints:

Project 16 should be completed before work on this project is initiated. Project 16 must resolve the question of future responsibility and will provide guidance concerning the degree of flooding to be attained.

Duration and Cost:

See Table 17.

B. PROJECT 10, DEVELOPMENT OF SURFACE WATER INFILTRATION CONTROL PROCEDURES

Objective:

To provide guidelines for minimizing the infiltration of surface water and design specifications for constructing surface-water diversion structures.

Work Tasks:

(1) Prepare a manual for use by the coal industry that provides guidelines for minimizing the creation of those conditions that result in infiltration of surface water. This manual should also contain design specifications for constructing surface-water diversion structures. The relative economics of water diversion control compared to the economics of pumping and treating acid mine water should be determined and included in the manual.

(2) A study should be undertaken to determine the need for and the minimum size requirements for pillars to provide adequate support for surface water bodies. These specifications are needed as current state requirements are so stringent that coal recovery is being minimized. For instance, currently proposed requirements in West Virginia will preclude the use of longwall mining at many mining sites.

Applicability:

Results of performing Task 1 will be applicable to all mines (see Table 15) where acid mine water may be generated and will require discharge to the surface.

Results of Task 2 will be applicable to all underground mining sites, whether or not acid water is formed, where the mine underlies any surface water body.

Duration and Cost:

See Table 17.

C. PROJECT 14, STUDY THE EFFECT OF ROOF COLLAPSE ON MINE WATER CHARACTERISTICS

Objective:

To determine if essentially complete roof collapse will result in the reduced flow of mine water and/or will inhibit the formation of acid mine water. Such collapse will essentially be complete following longwall mining or pillaring. In effect, this may be found to be an abatement measure if compared to abandoned mines where coal recovery was limited.

Work Tasks:

(1) A study should be undertaken to evaluate the effect of roof collapse on mine water quality and quantity. To conduct this study, no less than ten sets of mines or sections of mines should be compared. Each set of mines should be in the same seam and be generally similar except that caving has occurred at only one site.

(2) Studies should also be undertaken to determine the effect of injecting highly-reactive alkaline material, or other potential inhibitors, into an abandoned, caved section. Since the collapsed section no longer represents a dynamic system, long-term inhibition of acid formation might result. These studies should also be conducted in conjunction with the section sealing studies (Project 17).

(3) If roof collapse is found to have a positive effect on mine water characteristics, mining procedures should be developed to optimize its use as an abatement procedure.

Applicability:

As an "at-source" abatement procedure, positive results of performing this project would be applicable to all mines where acid water may be formed and become a polluttional problem. As a technique that relates to mine closure, successful results would be applicable to new mines in 25 percent of the coal reserves in Northern Appalachia and 10 percent of the coal reserves in Southern Appalachia.

Sequential Restraints:

Performance of Task 3 will depend upon an evaluation of the results obtained from performing Tasks 1 and 2.

Duration and Cost:

See Table 17.

D. PROJECT 16, EVALUATION OF THE LONG-TERM EFFECTS OF FLOODING UNDERGROUND MINES ABOVE DRAINAGE

Objective:

To determine under what conditions complete or partial mine flooding should be the objective of the mine-closure procedures. Flooding will be the result of mining on the dip or of construction of effective seals. Impounding of these vast quantities of water represents potential future hazards that must be considered when recommending that flooding be achieved. These hazards need to be more clearly defined and a program must be established by a responsible government agency to ensure that catastrophic events will not occur. Such events might include future construction projects that would break into a flooded mine, thereby releasing the impounded water, and possibly resulting in excessive stream pollution or even death and destruction.

Work Tasks:

(1) The Bureau of Mines should implement a continuing program to map those mines where flooding has occurred and as it occurs in the future. This program should include periodic monitoring of the sites to ensure that catastrophic events will not result from surface or underground activities that might interfere with the integrity of the system. If the Bureau or other government agency is unwilling to accept this responsibility, then flooding should be limited to small (to be defined by research) mines and/or mines with a relatively low (to be defined by research) hydrostatic head.

(2) Determine the effect of flooding on ground water quality and the general ground water hydrology.

(3) Establish a program to monitor several recently (within the last ten years) completed mine-sealing projects. Funds to provide continuous or even periodic monitoring of many of these sites are not currently available. Such a study is necessary to reliably evaluate the sealing procedures used to determine their long-term effectiveness in minimizing acid mine drainage pollution and their durability. Funds should also be made available for maintenance of the structures during the study period.

(4) Evaluate and define the potential dangers that would result from the accidental release of water impounded in a flooded mine. Such a study should define the activities that would cause the release and should consider restrictions on the hydrostatic head and volumes of water impounded.

Applicability:

Results will be applicable to new mines opened in coal seams that represent approximately 25 percent of the reserves in Northern Appalachia and 10 percent of the reserves in Southern Appalachia.

Duration and Cost:

See Table 17.

E. PROJECT 17. INVESTIGATE THE EFFECTIVENESS OF SECTION SEALING IN ACTIVE MINES

Objective:

To determine if section sealing is an effective "at-source" abatement technique.

Work Tasks:

(1) Evaluate the effects and cost effectiveness of partial or complete flooding of sections of mines on the quality and quantity of the mine water produced. Design criteria should be established for the construction of the various impoundment mechanisms used and the size of pillars that must be left in place. Such mechanisms would range from simple retention dams to hydraulic seals.

(2) Evaluate the effects of section air sealing on the quality of water produced. Since significant reduction in oxygen content has been shown to partially inhibit acid formation, this type of sealing might have application in deeper mines where the overburden is not fractured to the surface.

Applicability:

All mines in seams that produce acid mine water.

Sequential Restraints:

Work on developing design criteria should not begin until effectiveness of methodology is established. Variances to restrictions imposed by ventilation requirements must be obtained prior to initiation of all work.

Duration and Cost:

See Table 17.

It is also recommended that the Bureau actively pursue projects (5) Develop a Simple Model for Predicting Mine Water Quality and Quantity, (12) Prepare a Manual on Mine Drainage Abatement Procedures, and (13) Develop a Program to Disseminate Information on Mine Drainage Abatement to Coal Industry. These projects would not result in the development of "at-source" abatement techniques but they would be of immediate value to the industry in that wider application of available "at-source" techniques would result.

F. PROJECT 5, DEVELOP A SIMPLE MODEL FOR PREDICTING MINE WATER QUALITY AND QUANTITY

Objective:

To provide the coal industry with the technology that will enable them to fairly accurately forecast the quality and quantity of mine water that a planned mine will generate. As the technical and financial resources of most companies are limited, the model should be simple and should not require the use of a computer. Availability of the model will be cost effective as the size of required treatment plants can be more accurately determined and the need for planned "at-source" abatement procedures will be demonstrated.

Applicability:

All new underground mines during the planning stages.

Sequential Restraints:

Some difficulties will occur in demonstrating the reliability of such a model. If some degree of success cannot be obtained during the first two years of the project, it should be dropped.

Duration and Cost:

See Table 17.

G. PROJECT 12, PREPARE A MANUAL ON MINE DRAINAGE ABATEMENT PROCEDURES

Objective:

To make the entire industry aware of the availability of abatement procedures and to provide sufficient design criteria that will permit their implementation. Most large companies employ an adequate staff of environmental personnel who are knowledgeable of such procedures. This information is not currently available in a concise, easily-understood form and, as a result, abatement procedures have not been implemented by many of the smaller operators. Such a manual is necessary to enable the total industry to implement the presently-available technology and, through periodic updates, will provide information concerning new technology that will be developed by the Bureau and others.

Applicability:

All presently active and future mines that have potential acid mine drainage problems or will benefit from reduced pumping requirements.

Sequential Restraints:

The manual should be prepared upon completion of Project 10, Task 1, and should be updated as new or improved technology becomes available.

Duration and Cost:

See Table 17.

H. PROJECT 13, DEVELOP A PROGRAM TO DISSEMINATE INFORMATION ON MINE DRAINAGE ABATEMENT TO COAL INDUSTRY

Objective:

To establish an information-transfer program that will result in the implementation of "at-source" abatement. It is envisaged that such a program could be conducted by the Bureau's State Liaison Officers. Whatever the designated office, it should actively interface with the industry and should be in the position of providing complete information concerning all aspects of the mine-drainage problem within the specified state or district. Such a program must be established as it is recognized that availability of technology does not ensure implementation.

Applicability:

This program eventually should address mine-water problems other than acid and should not be limited to underground mines. It would then be of value to the entire industry.

Sequential Restraints:

If some type of information-transfer program is not established, it is doubtful if widespread implementation of "at-source" controls will be established. The overall value of the Bureau of Mines' involvement in mine drainage abatement would be significantly reduced and, in this respect, all proposed projects should be critically reviewed before being pursued.

Duration and Cost:

See Table 17.

TABLE 17. PROJECT WORK SCHEDULE AND ANTICIPATED MINIMUM COSTS

	1st Year	2nd Year	3rd Year	4th Year	5th Year	Additional Time Needed	Cost
6. Develop Mine Closure Technology							
1. Coal Barrier Specifications		██████████	██████████	██████████	██████████		200,000
2. Grouting Technology		██████████	██████████	██████████	██████████		500,000
3. Mine Seal Design		██████████	██████████	██████████	██████████		400,000
10. Develop Infiltration Controls							
1. Guidelines and Design	██████████						50,000
2. Pillar Size Specifications	██████████	██████████					75,000
14. Effect of Roof Collapse							
1. Effect on Mine Water Quality	██████████	██████████	██████████				500,000
2. Examine Use of Inhibitors		██████████	██████████	██████████			400,000
3. Mining Procedure Development				██████████	██████████		100,000
16. Evaluation of Flooding							
1. Maintenance and Monitoring	██████████	██████████	██████████	██████████	██████████	500 years	250,000/yr
2. Effect on Ground Water	██████████	██████████					200,000
3. Evaluate Completed Projects	██████████	██████████	██████████	██████████	██████████	10 years	250,000
4. Define Hazards	██████████						75,000
17. Investigate Section Sealing							
1. Evaluate Effects of Flooding	██████████	██████████	██████████				450,000
2. Evaluate Air Sealing		██████████	██████████	██████████	██████████		400,000
5. Develop Simple Predictive Model				██████████	██████████		200,000
12. Prepare Procedure Manual		██████████	██████████	██████████	██████████		50,000/update
13. Maintain Information Transfer Program						20 years	250,000/yr

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	701	729									
water infiltration											
	169	320	356	382	392	394	427	437	460	646	719
	741	745									
inhibitors											
biological											
	73	79	167	255	256	265	435	569	570	571	572
	625	649	669	670	671						
coatings											
	2	111	136	137	174	196	278	286	333	400	497
	519	577	655								
inert gas atmospheres											
	95	96	269	374	548	549	552	554	734		
in-situ neutralization											
	386	609									
mine closure											
	279	280	282	355	414	503	543				
air sealing											
	9	124	126	130	155	165	276	329	334	353	356
	364	365	393	395	413	415	416	418	432	438	461
	477	478	479	480	481	483	491	510	518	537	564
	617	641	642	722							
backfilling											
	517										
hydraulic sealing											
	123	128	270	271	272	273	283	284	499	534	563
	639	745									

Abatement methods											
mine closure (cont.)											
materials	169	172	270	271	273	382	443	499	523	524	609
mining methods	214	269	457	512	536	566					
mine barrier width	462	714	715	716							
preplanning	346										
Acid mine drainage formation	5	7	13	89	118	125	130	134	141	162	175
	325	327	330	331	332	351	354	366	424	425	503
	509	514	534	544	547	575	587	588	625	627	632
	633	703	772								
chemical oxidation	77	84	85	133	148	157	159	160	174	176	252
	294	321	324	486	487	488	489	490	491	496	504
	557	559	567	576	583	584	585	589	590	591	612
	613	725	759	761	770						
chemical versus biological acid formation	73	78	80	81	82	83	158	177	257	401	405
	426	434	528	556	578	586	592	614	658		
microorganisms in acid mine drainage	119	136	147	156	187	188	255	256	357	358	359
	360	361	362	363	402	404	406	407	408	433	435
	539	560	579	580	581	620	622	623	624	626	628
	629	630	631	634	635	636	648	649	669	672	682
	683	744	748	759							
Geology	76	142	211	213	217	218	343	657	772		
Alabama, Mary Lee coalbeds	495										
coalbed lithology	561	732	733	749							
Alabama	19	215	377	551							
Alaska	711	735	758	767	771						
Arkansas	21	511									
Colorado	22	241	242								
Illinois	23	290									
Indiana	24										
Iowa	25										

Geology

coalbed lithology (cont.)

Kansas	26										
Kentucky	27	204	206	207	384	466	468	665	666	667	
Maryland	203	645	668								
Missouri	30										
Montana	31	296									
New Mexico	32	242									
Ohio	33	146	153	707							
Oklahoma	34										
Pennsylvania											
anthracite	35	558	709								
underclays	371										
bituminous	36	72	109	110	201	202	205	208	209	210	310
	312	313	318	441	601	659	660	661	662	705	708
Tennessee	37	38	347	348	349	350	428	687	688	689	690
	691	692	693	694	695						
Utah	41										
Virginia	42	43	154	232	237	238	307	308	463		
Washington	44	45	699								
West Virginia	46	47	231	233	234	235	236	239	240	247	248
	249	306	309	311	314	315	316	317	336	337	469
	470	471	538	621	647	663	664	706	730		
western coals	20	29	297								
Wyoming	48	243	644	730							
conditions affecting mining and drainage formation	88	113	244	259	260	322	392	427	439	447	448
	449	450	451	452	456	482	508	527			
deposition and formation of coal	66	112	161	287	302	335	338	391	409	513	526
	542	555	562	594	615	637	673	674	678	680	686
	757										

Geology											
hydrology	13	85	92	490	510						
Indiana	12										
Kentucky	222	727	746	767							
Maryland	367	763									
Ohio	14										
Pennsylvania	69	70	91	93	144	186	261	277	445	458	616
West Virginia	768										
limestone occurrence	295	505									
mining effects on overlying strata	149	150	151								
stratigraphy	74	75	330	596	731	743					
Alabama	377										
Alaska	718										
Appalachia	66	212	262	602							
Arkansas	323										
Colorado	399	429									
Illinois	18	180	595	696							
Indiana	373	600									
Kentucky	372	387	467	468							
Maryland	645										
Ohio	99	138	139	140	223	227	228	229	304	681	
Pennsylvania	195	197	220	221	390	516	568	573	653	654	769
Virginia	143	472									
West Virginia	270	271	464	465	469	470	471				
western United States	739										

Geology											
stratigraphy (cont.)											
Wyoming	100										
Mining methods, effect on mine drainage formation	98										
Preparation, washability studies, and coal analyses											
	39	40	50	51	52	53	54	55	56	57	58
	59	60	61	62	63	64	65	163	224	299	300
	301	302	303	379	380	381	442	455	532	651	656
	713	731	732	733	743	749	762				
Alabama	19	293	377	525	551	574					
Alaska	711	718	735	758	767	771					
Appalachia	289	602									
Arkansas	21	292	511								
Colorado	22										
eastern coals	546	640									
Illinois	23	152	198	290	340	341	342	368	638		
Indiana	24	600									
Iowa	25	164	185								
Kansas	26										
Kentucky	27	204	206	207	384	466	467	468	665	666	667
Maryland	28	203	645	668							
Missouri	30										
Montana	31	296									
New Mexico	32										
Ohio	33	138	139	140	146	223	227	228	229	707	
Oklahoma	34										
Pennsylvania											
anthracite	35	116	375	558	709						

Geology											
stratigraphy (cont.)											
Wyoming											
	100										
Mining methods, effect on mine drainage formation											
	98										
Preparation, washability studies, and coal analyses											
	39	40	50	51	52	53	54	55	56	57	58
	59	60	61	62	63	64	65	163	224	299	300
	301	302	303	379	380	381	442	455	532	651	656
	713	731	732	733	743	749	762				
Alabama											
	19	293	377	525	551	574					
Alaska											
	711	718	735	758	767	771					
Appalachia											
	289	602									
Arkansas											
	21	292	511								
Colorado											
	22										
eastern coals											
	546	640									
Illinois											
	23	152	198	290	340	341	342	368	638		
Indiana											
	24	600									
Iowa											
	25	164	185								
Kansas											
	26										
Kentucky											
	27	204	206	207	384	466	467	468	665	666	667
Maryland											
	28	203	645	668							
Missouri											
	30										
Montana											
	31	296									
New Mexico											
	32										
Ohio											
	33	138	139	140	146	223	227	228	229	707	
Oklahoma											
	34										
Pennsylvania											
anthracite											
	35	116	375	558	709						

Preparation, washability studies, and coal analyses

Pennsylvania (cont.)											
bituminous											
	36	72	189	199	200	201	202	205	208	209	210
	288	310	312	313	318	441	516	597	601	653	659
	660	661	662	705	708						
Tennessee											
	37	38	347	348	349	350	428	687	688	689	691
	692	693	694	695							
Utah											
	41										
Virginia											
	42	43	232	237	238	307	308	463			
Washington											
	44	45	699								
West Virginia											
	46	47	231	233	234	235	236	239	240	306	309
	311	314	315	316	317	336	337	464	465	469	470
	471	538	621	647	663	664	706	730			
western coals											
	20	29	297	444	739						
Wyoming											
	48	644	730								
Pyrite occurrence											
	163	164	221	242	288	289	340	341	342	410	411
	446	656	757								
framboidal form											
	453	723									
Water quality data											
general											
	120										
Pennsylvania											
	103										
ground water quality in mining areas											
	86										
Illinois											
	595	737									
Kentucky											
	494	723									
Pennsylvania											
	158	162	291	459	500	501	529	530	531		
mine effluent quality											
	90	108	117	127	130	132	263	366	376	415	518
	611										
Colorado											
	750										
Indiana											
	418										
Maryland											
	367	370	475	476	679						

Water quality data
mine effluent quality (cont.)

Ohio	266	267	389	432	599	610					
Pennsylvania	4	11	17	87	97	98	145	179	191	192	219
	254	261	275	281	298	322	328	339	385	396	411
	413	417	421	436	445	474	498	506	507	568	598
	603	604	605	606	675	684	698	700	747	769	
West Virginia	171	194	270	344	345	356	378	383	420	454	492
	540	544	545	550	603						
stream quality in mining areas	101	121	135	253							
Alabama	742										
Appalachia	102	178	230	503	515	611					
Illinois	724										
Indiana	12	193									
Kentucky	494	727	738	767							
Maryland	369	679	726								
Missouri	721										
Ohio	181	266	610	717							
Pennsylvania	10	49	94	173	191	192	275	412	413	557	685
	701	702	720	747	756	770					
West Virginia	9	264	768								