

ACCELERATED DISTRIBUTION DEMONSTRATION SYSTEM

REGULATORY INFORMATION DISTRIBUTION SYSTEM (RIDS)

ACCESSION NBR: 8903090088 DOC. DATE: 89/02/24 NOTARIZED: YES DOCKET #
 FACIL: 50-315 Donald C. Cook Nuclear Power Plant, Unit 1, Indiana & 05000315
 50-316 Donald C. Cook Nuclear Power Plant, Unit 2, Indiana & 05000316
 AUTH. NAME AUTHOR AFFILIATION
 ALEXICH, M.P. Indiana Michigan Power Co. (formerly Indiana & Michigan Ele
 RECIP. NAME RECIPIENT AFFILIATION
 MURLEY, T.E. Document Control Branch (Document Control Desk)

SUBJECT: Responds to recommendations made in 880808 Generic Ltr 88-14
 re instrument air supply sys problems.

DISTRIBUTION CODE: A048D COPIES RECEIVED: LTR 1 ENCL 1 SIZE: 30
 TITLE: OR/Licensing Submittal: Equipment Qualification

NOTES:

RECIPIENT ID CODE/NAME	COPIES LTTR ENCL	RECIPIENT ID CODE/NAME	COPIES LTTR ENCL
PD3-1 LA	1 0	PD3-1 PD	1 0
STANG, J	1 1		
INTERNAL: ARM/DAF/LFMB	1 0	NRR/DEST/ADE 8H	1 0
NRR/DEST/MEB 9H	1 1	NRR/DEST/SGB 8D	1 1
NUDOCS-ABSTRACT	1 1	OGC/HDS1	1 1
REG FILE 01	1 1	RES/DSIR/EIB	1 1
EXTERNAL: LPDR	1 1	NRC PDR	1 1
NSIC	1 1		

NOTE TO ALL "RIDS" RECIPIENTS:

PLEASE HELP US TO REDUCE WASTE! CONTACT THE DOCUMENT CONTROL DESK,
 ROOM P1-37 (EXT. 20079) TO ELIMINATE YOUR NAME FROM DISTRIBUTION
 LISTS FOR DOCUMENTS YOU DON'T NEED!

TOTAL NUMBER OF COPIES REQUIRED: LTTR 14 ENCL 10

R
I
D
S
/
A
D
D
S



AEP:NRC:1075

Donald C. Cook Nuclear Plant Units 1 and 2
Docket Nos. 50-315 and 50-316
License Nos. DPR-58 and DPR-74
INSTRUMENT AIR SUPPLY SYSTEM PROBLEMS AFFECTING
SAFETY-RELATED EQUIPMENT

U.S. Nuclear Regulatory Commission
Document Control Desk
Washington, D. C. 20555

Attn: T. E. Murley

February 24, 1989

Dear Dr. Murley:

Generic Letter 88-14, "Instrument Air Supply System Problems Affecting Safety-Related Equipment," was issued on August 8, 1988. The Generic Letter follows up on an earlier NRC transmittal, NRC Information Notice 87-28 Supplement 1, which identified concerns related to the adverse effects on safety-related equipment from instrument air system failures. Generic Letter 88-14 was issued to request that each licensee review the earlier NRC transmittals and perform a design and operations verification of its instrument air systems. The purpose of this letter is to provide the response to Generic Letter 88-14 with respect to Donald C. Cook Nuclear Plant. Responses to each of the Generic Letter's recommendations are provided in the attachments to this letter.

Specifically, Attachment 1 provides a brief description of Cook Nuclear Plant's compressed air system (including the control/instrument air system) and the separate diesel generator air system. Attachment 2 is the response to Recommendation No. 1 involving the air quality of the control air system. Attachment 3 provides the response to Recommendation No. 2 that includes a description of system maintenance practices, emergency procedures and training. Finally, Attachment 4 addresses Recommendation No. 3 regarding verification of design and testing of the control air system.

A review of the attachments will demonstrate that emphasis has been placed on the quality and reliability of the air systems at the Cook Nuclear Plant. It is believed that the existing practices and improvements regarding the compressed air systems are responsive to the NRC's concerns as expressed in Generic Letter 88-14.

8903090088 890224
PDR ADOCK 05000315
P PDC

A046
11

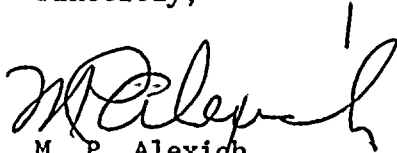
Dr. T. E. Murley

-2-

AEP:NRC:1075

This letter is submitted pursuant to 10 CFR 50.54(f) and, as such, an oath of affirmation is enclosed.

Sincerely,

A handwritten signature in cursive script, appearing to read 'M. P. Alexich', with a small vertical line above the 'h'.

M. P. Alexich
Vice President

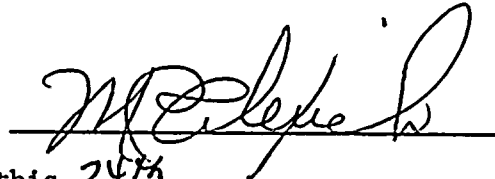
ldp

Attachments

cc: D. H. Williams, Jr.
W. G. Smith, Jr. - Bridgman
R. C. Callen
G. Charnoff
A. B. Davis
NRC Resident Inspector - Bridgman
G. Bruchmann

STATE OF OHIO)
COUNTY OF FRANKLIN)

Milton P. Alexich, being duly sworn, deposes and says that he is the Vice President of licensee Indiana Michigan Power Company, that he has read the foregoing Response to NRC Generic Letter 88-14, "Instrument Air Supply Problems Affecting Safety-Related Equipment," and knows the contents thereof; and that said contents are true to the best of his knowledge and belief.



Subscribed and sworn to before me this 24th

day of February, 1989.


NOTARY PUBLIC

BARBARA ANN BELVILLE
ATTORNEY AT LAW
NOTARY PUBLIC - STATE OF OHIO
MY COMMISSION HAS NO EXPIRATION DATE
SECTION 147.03 R. C.

1972

1972

1972



RECEIVED AND FORWARDED
BY THE SECRETARY OF STATE
ON 10/10/72
10/10/72
10/10/72

ATTACHMENT 1 TO AEP:NRC:1075

DESCRIPTION OF THE COMPRESSED AIR SYSTEM
AT THE COOK NUCLEAR PLANT

1. The first part of the document is a list of names and addresses of the members of the committee. The names are listed in alphabetical order, and the addresses are given in full, including the street, city, and state.

2. The second part of the document is a list of the names and addresses of the members of the committee who have been elected to the office of the secretary. The names are listed in alphabetical order, and the addresses are given in full, including the street, city, and state.

3. The third part of the document is a list of the names and addresses of the members of the committee who have been elected to the office of the treasurer. The names are listed in alphabetical order, and the addresses are given in full, including the street, city, and state.

4. The fourth part of the document is a list of the names and addresses of the members of the committee who have been elected to the office of the clerk. The names are listed in alphabetical order, and the addresses are given in full, including the street, city, and state.

I. COMPRESSED AIR SYSTEM (CAS)

A brief description of the compressed air system, which includes the control air system (also known as the instrument air system), is provided in this attachment for a better understanding of this response to Generic Letter 88-14. A simplified schematic diagram of the CAS for both units of the Cook Nuclear Plant is attached. In addition, the air system for the emergency diesel generators, which is a separate system from the plant's CAS, is described at the end of this attachment.

A. Design Bases

The intended functions of the CAS are:

1. The system must provide reliable compressed air supplies for control and instrument air requirements.
2. The system must provide adequate compressed air capacity for:
 - a. General Plant Service
 - b. Control
 - c. Instrumentation
 - d. Testing
 - e. Containment Penetration and Weld Channel Pressurization System
 - f. Respiratory protection in the containment per compressed gas association commodity Spec. G-7.1 - 1966 and OSHA Standards and Interpretations 1910.134.
3. The system should provide a continuous supply of compressed air to vital systems under both normal and abnormal conditions, although no credit is taken for the system in the Cook Nuclear Plant's Final Safety Analysis Report.

B. Component and System Design

1. Compressors and Associated Equipment

The CAS is designed to provide a reliable supply of compressed air for all plant uses. The CAS has two, oil-free, constant speed, 3-stage, motor-driven centrifugal plant air compressors (PACs) that provide about 1500 scfm of compressed air each at a discharge pressure of 100 psig at an inlet temperature of 90°F. During normal operation, only one PAC is in operation, supplying the demand for all plant air (including breathing air) and control/instrument air for the

entire plant. A 100% reserve capacity is, therefore, provided by the standby PAC. To prevent problems associated with moisture and oil in the CAS, the compressed air must be relatively moisture- and oil-free. In addition to using "oil-free" compressors, each compressor has been provided with an aftercooler, which includes a moisture separator. The aftercooler maintains the temperature of the compressed air that is supplied to the receivers below 95°F. The increased air moisture content, resulting from aftercooling, is removed by a cyclone-type separator. A plant air receiver is located downstream of each aftercooler. The air receivers accommodate sudden or unusually heavy demands for compressed air and prevent compressor surge.

The two "oil-free" control air compressors (CACs), one per unit, are constant speed, motor-driven, double acting, water cooled, vertical, reciprocating compressors that provide about 317 scfm of compressed air at a discharge pressure of 100 psig with an air inlet temperature of 70°F. Each CAC is also capable of supplying air to the containment penetration and weld channel pressurization system for its unit. Each CAC discharges through an aftercooler to a wet control air receiver. If both plant compressors are unavailable, low air pressure at each unit's wet control air receiver will automatically start both CACs. The CACs will run at a regulated, constant speed until stopped by the operator. The CAS has two parallel 100% capacity dry control air receivers and upstream strings of prefilters, air dryers, and after filters, thus assuring clean, dry air for instrument and control air usage. Each dryer consists of dual adsorption towers filled with a silica gel desiccant. The compressed air is dried as it passes through the desiccant bed of the adsorption tower. Two parallel 5-micron prefilters and two parallel 5-micron after filters are located before and after each dryer to prevent contamination of the dryer desiccant (moisture and pipe scale carry-over) and to protect the instruments and controls against desiccant dusting.

The control air system includes sufficient storage capacity to supply the control and instrument air requirements with the equivalent of approximately 5 minutes of control air output. Additionally, certain vital control valves within the containment are each equipped with a local receiver tank with enough capacity to activate the valve. Also, with operator action, all air compressors and their dryers can be supplied with electric power from normal, offsite and emergency sources.

[illegible][illegible]

1. *Introduction*
 2. *Background*
 3. *Methodology*
 4. *Results*
 5. *Discussion*
 6. *Conclusion*
 7. *References*
 8. *Appendix*
 9. *Tables*
 10. *Figures*
 11. *Supplementary Materials*
 12. *Notes*
 13. *Abbreviations*
 14. *Conflicts of Interest*
 15. *Acknowledgments*
 16. *Author Contributions*
 17. *Funding*
 18. *Data Availability Statement*
 19. *References*
 20. *Appendix*
 21. *Tables*
 22. *Figures*
 23. *Supplementary Materials*
 24. *Notes*
 25. *Abbreviations*
 26. *Conflicts of Interest*
 27. *Acknowledgments*
 28. *Author Contributions*
 29. *Funding*
 30. *Data Availability Statement*
 31. *References*
 32. *Appendix*
 33. *Tables*
 34. *Figures*
 35. *Supplementary Materials*
 36. *Notes*
 37. *Abbreviations*
 38. *Conflicts of Interest*
 39. *Acknowledgments*
 40. *Author Contributions*
 41. *Funding*
 42. *Data Availability Statement*
 43. *References*
 44. *Appendix*
 45. *Tables*
 46. *Figures*
 47. *Supplementary Materials*
 48. *Notes*
 49. *Abbreviations*
 50. *Conflicts of Interest*
 51. *Acknowledgments*
 52. *Author Contributions*
 53. *Funding*
 54. *Data Availability Statement*
 55. *References*
 56. *Appendix*
 57. *Tables*
 58. *Figures*
 59. *Supplementary Materials*
 60. *Notes*
 61. *Abbreviations*
 62. *Conflicts of Interest*
 63. *Acknowledgments*
 64. *Author Contributions*
 65. *Funding*
 66. *Data Availability Statement*
 67. *References*
 68. *Appendix*
 69. *Tables*
 70. *Figures*
 71. *Supplementary Materials*
 72. *Notes*
 73. *Abbreviations*
 74. *Conflicts of Interest*
 75. *Acknowledgments*
 76. *Author Contributions*
 77. *Funding*
 78. *Data Availability Statement*
 79. *References*
 80. *Appendix*
 81. *Tables*
 82. *Figures*
 83. *Supplementary Materials*
 84. *Notes*
 85. *Abbreviations*
 86. *Conflicts of Interest*
 87. *Acknowledgments*
 88. *Author Contributions*
 89. *Funding*
 90. *Data Availability Statement*
 91. *References*
 92. *Appendix*
 93. *Tables*
 94. *Figures*
 95. *Supplementary Materials*
 96. *Notes*
 97. *Abbreviations*
 98. *Conflicts of Interest*
 99. *Acknowledgments*
 100. *Author Contributions*
 101. *Funding*
 102. *Data Availability Statement*
 103. *References*
 104. *Appendix*
 105. *Tables*
 106. *Figures*
 107. *Supplementary Materials*
 108. *Notes*
 109. *Abbreviations*
 110. *Conflicts of Interest*
 111. *Acknowledgments*
 112. *Author Contributions*
 113. *Funding*
 114. *Data Availability Statement*
 115. *References*
 116. *Appendix*
 117. *Tables*
 118. *Figures*
 119. *Supplementary Materials*
 120. *Notes*
 121. *Abbreviations*
 122. *Conflicts of Interest*
 123. *Acknowledgments*
 124. *Author Contributions*
 125. *Funding*
 126. *Data Availability Statement*
 127. *References*
 128. *Appendix*
 129. *Tables*
 130. *Figures*
 131. *Supplementary Materials*
 132. *Notes*
 133. *Abbreviations*
 134. *Conflicts of Interest*
 135. *Acknowledgments*
 136. *Author Contributions*
 137. *Funding*
 138. *Data Availability Statement*
 139. *References*
 140. *Appendix*
 141. *Tables*
 142. *Figures*
 143. *Supplementary Materials*
 144. *Notes*
 145. *Abbreviations*
 146. *Conflicts of Interest*
 147. *Acknowledgments*
 148. *Author Contributions*
 149. *Funding*
 150. *Data Availability Statement*
 151. *References*
 152. *Appendix*
 153. *Tables*
 154. *Figures*
 155. *Supplementary Materials*
 156. *Notes*
 157. *Abbreviations*
 158. *Conflicts of Interest*
 159. *Acknowledgments*
 160. *Author Contributions*
 161. *Funding*
 162. *Data Availability Statement*
 163. *References*
 164. *Appendix*
 165. *Tables*
 166. *Figures*
 167. *Supplementary Materials*
 168. *Notes*
 169. *Abbreviations*
 170. *Conflicts of Interest*
 171. *Acknowledgments*
 172. *Author Contributions*
 173. *Funding*
 174. *Data Availability Statement*
 175. *References*
 176. *Appendix*
 177. *Tables*
 178. *Figures*
 179. *Supplementary Materials*
 180. *Notes*
 181. *Abbreviations*
 182. *Conflicts of Interest*
 183. *Acknowledgments*
 184. *Author Contributions*
 185. *Funding*
 186. *Data Availability Statement*
 187. *References*
 188. *Appendix*
 189. *Tables*
 190. *Figures*
 191. *Supplementary Materials*
 192. *Notes*
 193. *Abbreviations*
 194. *Conflicts of Interest*
 195. *Acknowledgments*
 196. *Author Contributions*
 197. *Funding*
 198. *Data Availability Statement*
 199. *References*
 200. *Appendix*
 201. *Tables*
 202. *Figures*
 203. *Supplementary Materials*
 204. *Notes*
 205. *Abbreviations*
 206. *Conflicts of Interest*
 207. *Acknowledgments*
 208. *Author Contributions*
 209. *Funding*
 210. *Data Availability Statement*
 211. *References*
 212. *Appendix*
 213. *Tables*
 214. *Figures*
 215. *Supplementary Materials*
 216. *Notes*
 217. *Abbreviations*
 218. *Conflicts of Interest*
 219. *Acknowledgments*
 220. *Author Contributions*
 221. *Funding*
 222. *Data Availability Statement*
 223. *References*
 224. *Appendix*
 225. *Tables*
 226. *Figures*
 227. *Supplementary Materials*
 228. *Notes*
 229. *Abbreviations*
 230. *Conflicts of Interest*
 231. *Acknowledgments*
 232. *Author Contributions*
 233. *Funding*
 234. *Data Availability Statement*
 235. *References*
 236. *Appendix*
 237. *Tables*
 238. *Figures*
 239. *Supplementary Materials*
 240. *Notes*
 241. *Abbreviations*
 242. *Conflicts of Interest*
 243. *Acknowledgments*
 244. *Author Contributions*
 245. *Funding*
 246. *Data Availability Statement*
 24

2. Compressed Air Distribution System

Compressed air is supplied to Units 1 and 2 through an air distribution system located in the turbine, auxiliary, and containment buildings. This distribution system consists of a shared plant air ring header extending throughout the turbine building, a pair of parallel plant air headers in the auxiliary building, and a plant and control air header in each containment. One PAC, plant air receiver, and plant air aftercooler is located in each unit. Each plant air receiver supplies its own plant air header in the auxiliary building; however, the shared turbine building plant air header may be supplied from either plant air receiver. The turbine building ring header, in turn, supplies compressed air to the standby plant air header in the auxiliary building. Since these distribution headers are parallel and have hose connections in essentially the same location, one unit's header may be isolated from the CAS without affecting the availability of plant air to the auxiliary building. Compressed air is then delivered to the header branches that feed the services requiring compressed air (e.g., screenhouse, service, turbine and auxiliary buildings).

Redundant containment isolation valves are provided on the instrument and control air headers going into both containments.

3. Piping, Valves and Pressure Vessel Design

All CAS equipment, piping, and valves are Seismic Class III with the exception of the Seismic Class I portion at the containment penetrations.

CAS equipment (e.g., receiver, dryer tanks, aftercoolers, and filter housings) has been designed and built in accordance with the ASME Boiler and Pressure Vessel Code, Section VIII criteria.

4. Cooling Water for Compressed Air System

The non-essential service water system (NESW) provides cooling water for the CAS. Two or three of the four pumps of the NESW system are normally operated to provide service water to the two units, with one or two pumps held in standby. All four pumps are able to take suction from either the Unit 1 or Unit 2 circulating water intake or discharge tunnels. Thus, if the tunnels of one unit are out of service, NESW supply to both units is ensured. Following a loss of all off-site power, the NESW pumps are automatically started after the emergency diesel generator power becomes available. Under these conditions, the pumps are primarily used to supply cooling water to the CACs in order to restore

1. The first part of the report is a general introduction to the subject of the study. It discusses the importance of the study and the objectives of the research. It also provides a brief overview of the methodology used in the study.

2. The second part of the report is a detailed description of the study area. It includes information about the location of the study area, the population of the study area, and the characteristics of the study area. It also discusses the data sources used in the study.

3. The third part of the report is a detailed description of the study results. It includes information about the findings of the study, the conclusions drawn from the findings, and the implications of the findings. It also discusses the limitations of the study and the need for further research.

4. The fourth part of the report is a detailed description of the study conclusions. It includes information about the overall findings of the study, the conclusions drawn from the findings, and the implications of the findings. It also discusses the limitations of the study and the need for further research.

5. The fifth part of the report is a detailed description of the study recommendations. It includes information about the recommendations made by the study, the reasons for the recommendations, and the implications of the recommendations. It also discusses the limitations of the study and the need for further research.

control and instrument air service. All motor-operated valves on the NESW systems are operated from the station battery system. Cross-ties between the pumps permit any one pump to supply the initial blackout requirements for both units. The flexibility of design indicated above ensures a ready supply of cooling water to the Control Air System components.

C. Air Quality

The plant air quality complies with the OSHA standards for respiratory use. Compressed air for respiratory use inside containment is provided within the containment pipe annulus. At this location, fittings are provided for connecting air masks to the plant air system.

D. Maintenance and Testing

The plant also has an active inservice testing (IST) program under which safety-related valves are periodically tested. The portion of the IST program relevant to GL 88-14 is discussed later in this submittal. Safety-related, air-operated valves are only used at the Cook Nuclear Plant when the failure mode is also the desired post-accident condition.

II. EMERGENCY DIESEL GENERATOR STARTING AIR SYSTEM

A. Design Bases

The intended functions of the starting air system (SAS) for the emergency diesel generators (EDG) are:

1. The system must provide redundant compressed air for starting the diesel engine of the EDG.
2. The system must provide adequate compressed air capacity for:
 - a. The engine of the EDG to start, accelerate up to speed, and be ready to accept EDG load within ten seconds from the initiation of the start signal
 - b. Other compressed air needs, i.e., control air and power for pneumatically operated valves in the system and for the flywheel air jack (turning gear).

B. Component and System Design

1. Compressors and Associated Equipment and Air Distribution System

Each engine has its own SAS that is totally independent of the plant CAS. Each system consists of two redundant air supply trains having a compressor, receiver, starting air supply valve and controls. Each supply train is capable of starting the engine. The two trains are cross-tied at the engine starting air manifolds. There is a check valve at the end of each train just upstream of the cross-tie line to prevent backflow should one supply train fail. Each train supplies air through an air-operated starting supply valve and a check valve. Additionally, an air-operated turbo jet assist valve enables the "fast" start of each engine on demand. The air supply to the turbo jet assist valves is through pressure regulators. The jet assist valves automatically open for the fast start.

As stated earlier, two starting air supply trains serve each engine. Each train consists of an air receiver to store the pressurized starting air and a motor-driven air compressor to maintain the air receiver pressure within a range of 220 and 240 psig. There is a cross-tie containing a manual valve between the two compressor discharge pipes. This valve is normally closed to isolate the two starting air supply trains. If, however, one of the air compressors is out of service, the valve will be opened by operator action to keep both air receivers properly pressurized.

The above redundancies in the compressors and their associated equipment as well as the supply and distribution lines of the SAS ensure adequate starting air for the EDGs.

Finally, since the emergency diesel generators are normally in a standby mode, the supporting starting air system compressors are operated only to maintain the associated air receivers charged to full pressure to ensure a fast starting capability. Given this intermittent operation, system air quality testing is not appropriate and is, therefore, not addressed in Attachment 2. Further, since diesel generator operability is addressed in the technical specification surveillance requirements, the maintenance practices for the dryers and compressors are included in the plant's Preventive Maintenance Program. They are, therefore, not described further in Attachment 3. The starting air supply valves and turbo jet assist valves described above are further discussed in Section C.5 of Attachment 4.

2. Piping, Valves and Pressure Vessel Design

The equipment and instrumentation in the starting air system is supplied by the diesel engine manufacturer, and is arranged and maintained in accordance with the manufacturer's recommendations.

The system equipment and piping are Seismic Class I. The exceptions are (1) the air compressors and their discharge piping to the first check valves, and (2) the line from the air jack to its root valve, which are Seismic Class III.

3. Electrical Power Supply

The only motors requiring electrical power in the diesel engine starting air system are the two air compressors serving each engine. The compressors are not needed during an engine start-up.

The several solenoid valves within the Worthington engine control module receive their power from the 250-volt DC circuit that supplies the diesel engine control bus. This 250-volt DC circuit remains energized during a blackout. The solenoid valves control the air signals to the starting air supply valves, jet-assist valve, and throttle shutdown cylinder. These air signals control the starting and stopping of the engine.

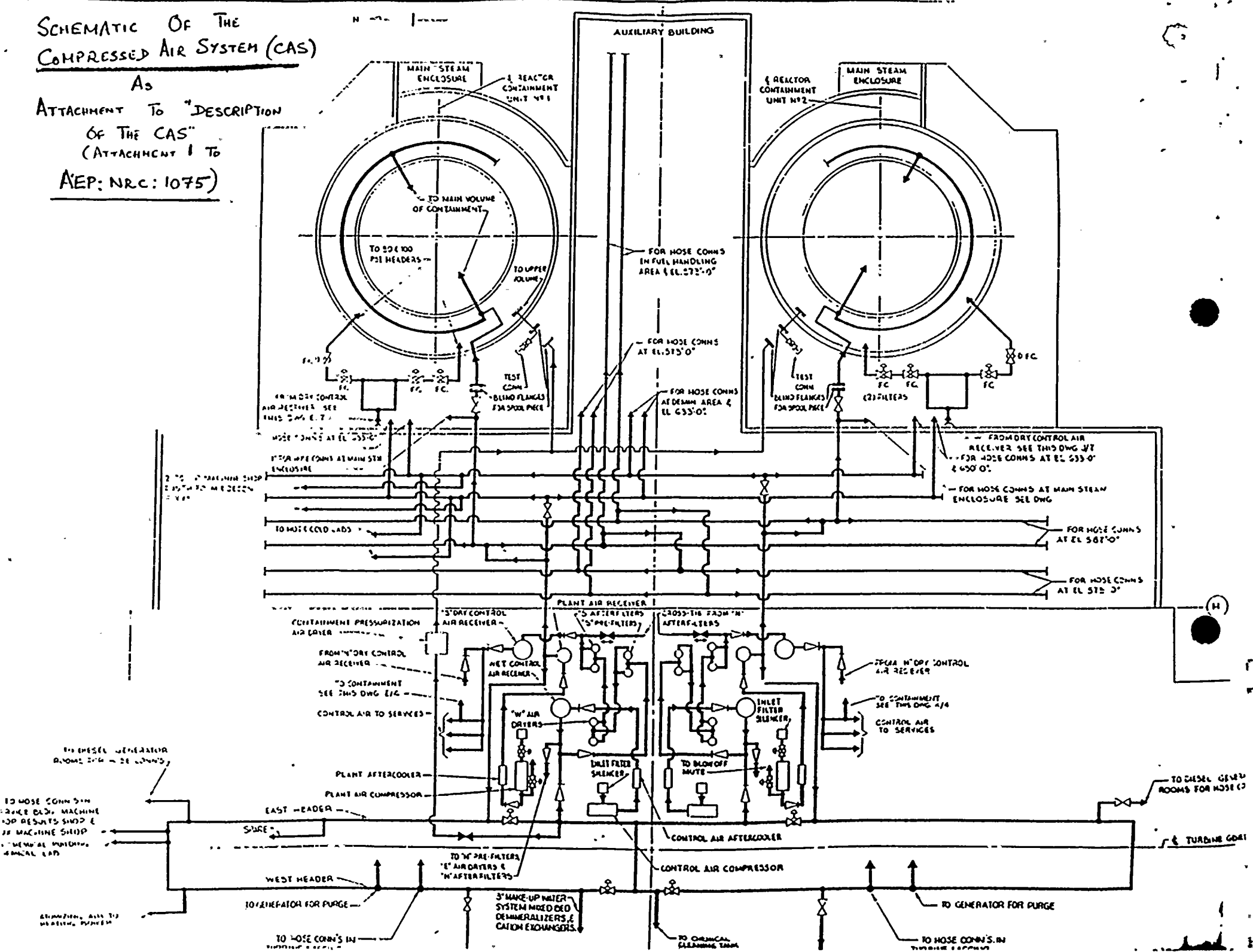


14-10

14-10

N 2. |

ATTACHMENT TO "DESCRIPTION
OF THE CAS"
(ATTACHMENT I TO
AEP: NRC: 1075)



ATTACHMENT 2 TO AEP:NRC:1075

RESPONSE TO RECOMMENDATION #1
OF GENERIC LETTER 88-14

I. RECOMMENDATION #1 - "Verification by test that actual instrument air quality is consistent with the manufacturer's recommendations for the individual components served."

II. RESPONSE

Summary

Manufacturers assume that their equipment will be provided with instrument quality air (in accordance with ANSI/ISA S 7.3-1975). Periodic sampling for dewpoint and hydrocarbons, combined with the filter arrangement and periodic delta-P monitoring in the plant and control air systems ensure that air of the proper quality is being supplied for safety-related equipment, as well as for important balance-of-plant equipment. Consistent with ANSI/ISA S 7.3-1975 and associated Regulatory Guide 1.68.3, elements of the air quality program include:

- i) maximum allowable moisture content,
- ii) maximum entrained particle size,
- iii) maximum allowable oil content,
- iv) awareness of possible corrosive or toxic contamination entering the air system.

A. Dewpoint

ANSI/ISA S 7.3 allows the dewpoint to be as high as 18°F below the minimum temperature to which any part of the instrument air system is exposed at any time of the year. Presently, air moisture is sampled at the Cook Nuclear Plant at the dry control air receiver tanks weekly. The dewpoint values have been satisfactory with respect to ANSI/ISA S 7.3, based on performance and test records available at the Cook Nuclear Plant.

In addition, the existing "Air Dryer Performance Test Procedure" uses the moisture content in the air at rated flow and pressure to determine equipment status, maintenance needs and operating efficiency of the instrument/control drying system. A modified program, which has been under development since the summer of 1988 and informally implemented, has been successful in identifying air dryer components susceptible to failure. This "Dryer Monitoring Program," which includes regularly scheduled replacement of certain components and the weekly dewpoint monitoring, will result in planned maintenance on the dryers rather than reactive maintenance. The Air Dryer Performance Test Procedure will be revised to become the Dryer Performance Monitoring Program, thereby moving the emphasis away from "acceptance" testing and towards enhancement of reliability.

ber

B. Particle Size

ANSI/ISA-S 7.3 specifies that "the maximum particle size in the air stream at the instrument shall be three (3) micrometers (microns)." The two trains of control air dryer after-filters at the Cook Nuclear Plant have a rated particle retention size of five (5) microns. There is no filter bypass. Additionally, pressure reducing station filters located downstream of the after-filters are rated as "removing 98% of all particulate matter greater than 0.07 microns." The control air to containment does not pass through this additional set of filters. An engineering analysis comparing instrument orifice sizes and dryer post-filter sizes was, however, completed in response to INPO SOER 88-01. This determined that the five-micron rating of the after-filters was adequate for protecting air quality for the air-operated equipment inside containment.

Particulate sampling is not currently being performed. Based on an engineering review of the maintenance history at Cook Nuclear Plant, documented in response to NRC IE IN 87-28, particulate sampling is not considered to be warranted at this time. Monthly differential pressure (ΔP) measurements are, however, being taken across the dryer after-filters as part of the Preventive Maintenance (PM) Program. Higher than normal readings would be indicative of unusual particulate or other loading of these filters. This program calls for the periodic monitoring and changeout of filter elements (including those in the pressure reducing stations) on a scheduled basis, before any significant deterioration can occur.

Pre- or after-filters would be replaced if filter ΔP as found during monthly testing approaches 12" H_2O . The plant has not observed/recorded any evidence that the ΔP across these filters had ever approached 12" H_2O . The dryer desiccant charge is replaced on an annual basis. This preventive maintenance is considered adequate to prevent particulate carryover into the CAS, based on manufacturers' recommendations.

By way of specific operating history, the after-filters were replaced in April 1988 as part of the PM program, and ΔP monitoring was instituted to track the "loading" of the dryers and filters. The differentials started out at, and remained at, between 1 and 2 inches of water, with no apparent loading of the filters after 8 months of operation.

Future operating history and test data of filter loading will be used to adjust the current monitoring program, as necessary.

The Cook Nuclear Plant dryers are of the down-flow design, which tends to compact the desiccant rather than keep it in suspension. This design feature results in minimizing desiccant dusting from the dryers and subsequent loading of the after-filters. The plant site has not observed/recorded any evidence of contamination of the control air system from desiccant carryover.

C. Oil/Hydrocarbon Content

Both sources of instrument/control air (the 2 centrifugal plant air compressors and the 2 reciprocating control air compressors - one for each unit) are of the "oil-free" type. The discharge of the plant air compressors is sampled on a monthly basis. This is performed in accordance with an established plant procedure to verify that the plant air system meets the requirements of OSHA CG A Type 1, Class D, "Breathing Air Supply." Analysis is performed by an offsite laboratory. The maximum allowable combined oil, mist, and hydrocarbon level under this Standard is 5mg/m³.

The maximum level of oil content allowable in the instrument/control air system per ANSI/ISA S 7.3 is 1 ppm (w/w). The composite oil, mist and hydrocarbon content of the plant air₃ has been analyzed to date as typically less than 0.2 mg/m³ (\approx 0.16 ppm w/w). Since control air is drawn from the plant air system without the introduction of additional contaminants, it follows that the oil content in the control air stream also complies with the Standard. The plant's sampling procedure requires notification of the Plant Technical Engineering Department if the 5 mg/m³ limit for breathing air is exceeded, so that the necessary corrective actions could be taken. No evidence exists of this limit having been exceeded.

D. Corrosive or Toxic Contamination

The air compressor intakes are located in the turbine building. Test results confirm that significant levels of contaminants are not entering the air stream.

THE UNIVERSITY OF CHICAGO
LIBRARY

ATTACHMENT 3 TO AEP:NRC:1075

RESPONSE TO RECOMMENDATION #2
OF GENERIC LETTER 88-14

THE
FEDERAL BUREAU OF INVESTIGATION
UNITED STATES DEPARTMENT OF JUSTICE

WASHINGTON, D. C.

TO : DIRECTOR, FBI
FROM : SAC, NEW YORK
SUBJECT: [Illegible]

RE: [Illegible]
[Illegible]
[Illegible]

DATE: [Illegible]
BY: [Illegible]

1. [Illegible]
2. [Illegible]
3. [Illegible]

4. [Illegible]
5. [Illegible]
6. [Illegible]

- I. RECOMMENDATION #2 - "Verification that maintenance practices, emergency procedures and training are adequate to ensure that safety-related equipment will function as intended on loss of instrument air."

II. RESPONSE

Summary

The maintenance practices, emergency procedures and training at the Cook Nuclear Plant are considered adequate to ensure that safety-related equipment will function as intended on loss of instrument or control air.

A. Maintenance Practices

Maintenance and post-maintenance activities must be adequate to ensure that safety-related equipment will function as required on loss of air. This requires that (1) maintenance on safety-related equipment and components be controlled, and (2) post-maintenance testing verify appropriate failure modes when considered necessary.

To satisfy these requirements at the Cook Nuclear Plant, a Job Order or approved instruction/procedure is required for all work on safety-related or safety-interface components. A Job Order is initiated whenever corrective maintenance (repair) is necessary for equipment, as well as when initiating modifications to equipment. A supervisory review by the department initiating the Job Order is also required to determine if the activity, or affected equipment, is Technical Specification (T/S) related. If so, the Job Order is appropriately marked to document the T/S impact and required post-maintenance testing. This information is again reviewed in the department assigned to accomplish the repairs or modification. Completion of required post-maintenance testing is also documented and attached to the Job Order itself.

Post-maintenance/modification testing requires the coordination of several groups, generally Operations (OPS), Instrument & Control (I&C), and Maintenance personnel. For situations dealing with air-operated, safety-related valves, Cook Nuclear Plant's operators use a special "Operations Standing Order" (OSO) that delineates actions to ensure proper inservice inspection timing for valves, both for routine and post-maintenance testing. This OSO also delineates specific examples of work after which post-maintenance testing is required. The Cook Nuclear Plant I&C Department's written policy is that "Post-Maintenance Testing shall be used to verify that equipment is repaired, operates correctly, and performs its desired function." The

policy also addresses temporary and permanent modifications and requires that post-modification testing be performed depending on the work/adjustments made. A design change implementation also often entails post-installation testing of safety-related equipment and components. The existing design process identifies all test requirements in accordance with a control valve specification for the fail-safe testing after a design change is performed. Routine fail-safe testing is performed in accordance with appropriate surveillance procedures. These procedures are derived from the Technical Specifications or the second ten-year Inservice Testing Program submitted to the NRC in 1987.

B. Emergency Procedures

Expected system and plant responses to a loss of instrument air are provided in the Cook Nuclear Plant emergency operating procedures for loss of control air and annunciator response procedures for plant systems. These emergency procedures include both automatic and required manual responses expected upon a reduction in control air system pressure. In addition, operating procedures for each unit specifically address responses to a loss of control air. These procedures list symptoms of air supply loss and identify the failure positions of key air-operated valves upon a complete air loss. In addition, the procedures direct operator action to trip the unit and begin a cooldown when the 100 psi control air header decreases to 80 psig. Subsequent operations are in accordance with the established emergency procedures for "Reactor Trip or Safety Injection" and "Reactor Trip Response" as applicable.

The combination of the existing emergency and operating procedures is considered to adequately address GL 88-14 in relation to the functioning of safety-related equipment on loss of instrument air.

C. Training

Emphasis on the importance of the instrument air system has been included in training for Maintenance (Mechanical Maintenance) personnel by including the recommendations of INPO's SOER 88-01 in a related lesson on Plant Air Systems. The importance is also stressed in the system training lesson plan for I&C personnel, which also includes specific problems of air systems mentioned in SOER 88-01 and NRC IN 87-28.

Operator training (utilizing INPO SOER 88-01 and NRC IN 87-28) has sensitized the Cook Nuclear Plant's operators to the importance of air systems and illustrated problems caused when air systems are contaminated or neglected. This training also serves to familiarize the operators with the

expected plant responses on a loss of air. Non-licensed plant operators, operators preparing to take an NRC licensing exam and licensed operators in the requalification program participated in classroom training in 1988 on loss-of-air. Simulator training on a loss-of-air scenario is planned for 1989.

ATTACHMENT 4 TO AEP:NRC:1075

RESPONSE TO RECOMMENDATION #3
OF GENERIC LETTER 88-14

116.
117.

- I. RECOMMENDATION #3: "Verification that the design of the entire instrument air system including air or other pneumatic accumulators is in accordance with its intended function, including verification by test that air-operated safety-related components will perform as expected in accordance with all design-basis events, including a loss of the normal instrument air system. This design verification should include an analysis of current air-operated component failure positions to verify that they are correct for ensuring required safety functions.

II. RESPONSE

Summary

The functions and design of the CAS and the emergency diesel generator starting air system at the Cook Nuclear Plant are described in Attachment 1 to the letter. The operating history of systems as well as the results of inservice or surveillance testing, indicate that the systems and their individual components can be expected to perform their intended safety-related functions. Further, the safety-related, air-operated valve failure positions have been verified to be correct during development of the Inservice Testing Program. The description of testing practices represent current commitments and may be changed to address future changes to the requirements without changes to this submittal in response to NRC GL 88-14.

A. Function of Air Systems

The CAS is designed with flexibility and reliability in mind, but it is not essential for the fulfillment of any safety function described in the Updated Safety Analysis Report. The functions of the CAS and the emergency diesel generator startup air system are described in Attachment 1.

B. Design of Air Systems

The design of the CAS and the SAS are described in Attachment 1. As indicated therein, the systems are capable of fulfilling their intended design functions. The SAS design was initially verified by pre-operational testing. Operability is demonstrated by the monthly running of the diesel generators in accordance with Technical Specification requirements.

The CAS has also performed well with no significant events attributable to failure of the control air system. The design and operation of "oil-free" compressors, filters, dryers, and receivers has contributed to a reliable air supply system. The use of welded stainless steel piping and

1. The first part of the document discusses the importance of maintaining accurate records of all transactions and activities. It emphasizes that proper record-keeping is essential for transparency and accountability, particularly in financial matters. The text suggests that organizations should implement robust systems to track every aspect of their operations, from procurement to sales, to ensure that all data is reliable and accessible.

2. The second part of the document addresses the challenges of data management in a rapidly changing environment. It highlights the need for continuous monitoring and updates to data systems to reflect the latest information. The author notes that while technology offers powerful tools for data collection and analysis, it also introduces complexities that require careful management. Organizations must therefore invest in training and resources to effectively utilize these tools and maintain the integrity of their data.

3. The third part of the document focuses on the importance of communication and collaboration between different departments and stakeholders. It argues that effective communication is key to ensuring that all parties are aligned and working towards common goals. The text suggests that regular meetings and open lines of communication can help to identify potential issues early on and facilitate the development of effective solutions. Collaboration is also emphasized as a critical factor in the success of any project or initiative.

4. The fourth part of the document discusses the role of leadership in driving organizational success. It stresses that leaders must be able to inspire and motivate their teams, as well as make strategic decisions that guide the organization's future. The text suggests that leaders should focus on creating a positive and supportive work environment where team members feel valued and empowered. Additionally, leaders should be proactive in identifying and addressing challenges, ensuring that the organization remains agile and responsive to change.

5. The fifth part of the document concludes by summarizing the key points discussed and offering final thoughts on the importance of maintaining high standards of performance and integrity. It reiterates that success is achieved through a combination of effective management, strong communication, and a commitment to excellence. The author encourages organizations to continue to strive for improvement and innovation, ensuring that they remain competitive in a dynamic market.

brazed copper tubing in the instrument and control air system has also reduced the risk of corrosion products adversely affecting the air system downstream of the filters.

C. Design Verification Testing

This section identifies the types of testing performed on the CAS and the emergency diesel generator starting air system for the purpose of verifying that the systems can perform as intended. Included is a description of the pre-operational testing, preventive maintenance, fail-safe testing and inservice testing programs as applicable to these systems.

1. Pre-operational Testing

Cook Nuclear Plant performed rapid and slow bleed down tests on the CAS in January 1975 and December 1977 on Units 1 and 2, respectively, with the results meeting the requirements of U.S. NRC Regulatory Guide 1.80. Testing was conducted in accordance with the Plant Loss of Instrument Air Pre-Operational Test Procedures.

2. Preventive Maintenance Program

The PACs, CACs, control air dryers, pre- and after-filters, and individual control air header filters are included in the plant's Preventive Maintenance Program. The program identifies the one- and five-year inspection requirements of the equipment as well as the relevant maintenance procedure. The one-year maintenance requirements are based on the manufacturer's technical manual and include 17 distinctive steps for inspection of the various parts and associated control functions of the CAC and 26 steps for the PAC. The five-year inspection/maintenance requires complete disassembly and inspection of components. The most recent inspection/preventive maintenance performed on the various CAS equipment is as indicated below.

	<u>Air Compressors</u>				<u>Control Air Dryer System</u>		
	<u>Annually</u>		<u>Five Year</u>		<u>Monthly</u>	<u>Annually</u>	
	<u>CAC</u>	<u>PAC</u>	<u>CAC</u>	<u>PAC</u>	<u>Suction Filter</u>	<u>Pre- and After-Filters</u>	<u>Individual Header Filters</u>
U1 10/21/88	3/18/88	2/4/87	11/18/86	1/24/89	4/12/88	4/12/88	
U2 10/28/88	1/21/89	12/11/87	12/9/87	1/19/89	5/4/88	5/4/88	

[illegible]

Figure 1. The effect of the concentration of the *Agrobacterium* suspension on the transformation efficiency of *Agrobacterium* strains. The *Agrobacterium* strains were grown in the YEA medium for 24 h at 28 °C. The cell concentration of the strains was adjusted to 1.0 × 10⁸ cells/ml. The cell suspension was mixed with the plant tissue and the transformation efficiency was determined. The results were expressed as the mean ± SD of three independent experiments. The asterisks indicate the significant difference between the strains at the same concentration of the cell suspension.

1. *Pharmaceutical industry* – The pharmaceutical industry is a major player in the healthcare sector, responsible for the development, production, and distribution of drugs. It is characterized by high R&D costs, long development cycles, and significant regulatory hurdles. The industry is often criticized for high prices and lack of transparency.

2. *Healthcare providers* – These include hospitals, clinics, and individual practitioners who deliver medical services. They are the primary point of contact for patients and are responsible for the diagnosis, treatment, and management of diseases.

3. *Insurance companies* – Insurance companies play a crucial role in financing healthcare. They collect premiums from individuals and businesses and use the funds to pay for medical services. They often negotiate with providers and pharmaceutical companies to secure lower rates.

4. *Government* – The government is involved in healthcare through regulation, funding, and ownership. It sets standards for safety and efficacy, provides funding for public health programs, and owns certain facilities, such as Veterans Affairs hospitals.

5. *Patients* – Patients are the end-users of healthcare services. They are responsible for seeking medical care, following treatment plans, and paying for services. Patient satisfaction and health outcomes are key concerns for all other stakeholders.

6. *Pharmaceutical distributors* – These companies act as intermediaries between manufacturers and healthcare providers. They manage the logistics of drug distribution, ensuring that medications are delivered to the right place at the right time.

7. *Medical device manufacturers* – These companies produce equipment and instruments used in medical procedures. They face similar challenges to pharmaceutical companies, including high R&D costs and regulatory requirements.

8. *Healthcare technology (HIT) companies* – These companies develop and provide software and hardware solutions for healthcare. Examples include electronic health record (EHR) systems, medical imaging equipment, and telemedicine platforms.

9. *Academic institutions* – Universities and research centers are involved in the discovery and development of new drugs and medical technologies. They often collaborate with the pharmaceutical industry and receive funding from government and private sources.

10. *Pharmaceutical associations* – These organizations represent the interests of pharmaceutical companies. They lobby on behalf of the industry, provide information to the public, and coordinate industry efforts.

11. *Healthcare reform advocates* – These groups advocate for changes to the healthcare system, such as universal coverage, cost containment, and improved access. They often work to influence public opinion and policy.

12. *Pharmaceutical regulators* – These agencies, such as the FDA in the US, are responsible for ensuring the safety and efficacy of drugs. They review applications for new drugs, monitor safety, and enforce regulations.

13. *Pharmaceutical wholesalers* – These companies purchase drugs in bulk from manufacturers and sell them to healthcare providers. They play a key role in the distribution chain.

14. *Pharmaceutical retailers* – These are the places where patients purchase their medications, such as pharmacies. They are responsible for dispensing drugs and providing patient education.

15. *Pharmaceutical manufacturers* – These are the companies that actually produce the drugs. They are responsible for the quality and safety of the manufacturing process.

16. *Pharmaceutical sales representatives* – These individuals are responsible for promoting and selling pharmaceutical products to healthcare providers. They often visit providers in person to discuss new products and provide samples.

17. *Pharmaceutical research and development (R&D)* – This is the process of discovering and developing new drugs. It involves a long and costly process of identifying potential targets, designing and synthesizing compounds, and testing them in preclinical and clinical studies.

18. *Pharmaceutical marketing* – This involves the promotion and sale of pharmaceutical products. It includes advertising, sales, and other promotional activities.

19. *Pharmaceutical distribution* – This is the process of getting drugs from the manufacturer to the patient. It involves a complex network of intermediaries, including wholesalers, distributors, and retailers.

20. *Pharmaceutical pricing* – This is the process of determining the price of a drug. It is a complex process that involves many factors, including R&D costs, manufacturing costs, and market competition.

21. *Pharmaceutical quality control* – This is the process of ensuring that drugs are manufactured to the highest standards of quality. It involves testing and monitoring at every stage of the manufacturing process.

22. *Pharmaceutical safety* – This is the process of ensuring that drugs are safe for use. It involves monitoring for adverse effects and taking action to remove unsafe products from the market.

23. *Pharmaceutical innovation* – This is the process of developing new drugs and medical technologies. It is a key driver of progress in healthcare and is often the focus of government and industry funding.

24. *Pharmaceutical regulation* – This is the process of creating and enforcing rules that govern the pharmaceutical industry. It is designed to protect public health and ensure the integrity of the healthcare system.

25. *Pharmaceutical transparency* – This is the process of making information about the pharmaceutical industry more accessible to the public. It includes disclosing financial relationships, clinical trial results, and other information that is relevant to patient care.

26. *Pharmaceutical access* – This is the process of ensuring that all patients have access to the care and services they need. It involves addressing barriers to care, such as cost, location, and availability.

27. *Pharmaceutical equity* – This is the process of ensuring that healthcare is distributed fairly and that all patients have an equal opportunity to benefit from medical advances.

28. *Pharmaceutical sustainability* – This is the process of ensuring that the healthcare system is able to meet the needs of future generations. It involves addressing issues such as climate change, resource scarcity, and social inequality.

29. *Pharmaceutical ethics* – This is the process of applying moral principles to the pharmaceutical industry. It involves addressing issues such as conflicts of interest, patient autonomy, and the distribution of resources.

30. *Pharmaceutical governance* – This is the process of overseeing the pharmaceutical industry and ensuring that it operates in a responsible and transparent manner. It involves setting standards, monitoring performance, and taking action when necessary.

31. *Pharmaceutical accountability* – This is the process of holding individuals and organizations in the pharmaceutical industry responsible for their actions. It involves establishing clear lines of responsibility and ensuring that those responsible are held accountable.

32. *Pharmaceutical integrity* – This is the process of ensuring that the pharmaceutical industry operates with honesty and transparency. It involves avoiding conflicts of interest and ensuring that all information is disclosed accurately.

33. *Pharmaceutical trust* – This is the process of building confidence in the pharmaceutical industry. It involves demonstrating a commitment to patient care, transparency, and ethical behavior.

34. *Pharmaceutical reputation* – This is the process of managing the public perception of the pharmaceutical industry. It involves addressing negative publicity and promoting positive stories that highlight the industry's contributions to healthcare.

35. *Pharmaceutical brand* – This is the name and identity of a pharmaceutical company or product. It is a key asset for the company and is used to distinguish its products from those of its competitors.

36. *Pharmaceutical intellectual property* – This is the legal right to a new drug or medical technology. It is granted to the inventor and allows them to control the use and distribution of the invention for a certain period of time.

37. *Pharmaceutical patent* – This is a legal document that grants the inventor of a new drug or medical technology the exclusive right to make, use, and sell the invention for a certain period of time.

38. *Pharmaceutical trademark* – This is a symbol, word, or design that identifies a particular product or company. It is used to distinguish the product from others in the market.

39. *Pharmaceutical copyright* – This is the legal right to a creative work, such as a book, movie, or song. It is granted to the creator and allows them to control the use and distribution of the work.

40. *Pharmaceutical trade secret* – This is information that is kept secret and provides a competitive advantage to a company. It is protected by law and can be a valuable asset for the company.

41. *Pharmaceutical litigation* – This is the process of resolving legal disputes between pharmaceutical companies or between a company and a patient. It can involve lawsuits, arbitration, or other legal proceedings.

42. *Pharmaceutical regulation* – This is the process of creating and enforcing rules that govern the pharmaceutical industry. It is designed to protect public health and ensure the integrity of the healthcare system.

43. *Pharmaceutical compliance* – This is the process of ensuring that a company follows all applicable laws and regulations. It involves establishing internal controls and monitoring compliance with external requirements.

44. *Pharmaceutical risk management* – This is the process of identifying and managing risks that could harm a company or its patients. It involves assessing the likelihood and impact of risks and taking steps to minimize them.

45. *Pharmaceutical crisis management* – This is the process of responding to a crisis, such as a product recall or a safety issue. It involves coordinating communication, managing the situation, and restoring trust.

46. *Pharmaceutical public relations* – This is the process of managing the relationship between a company and the public. It involves creating and disseminating information that is accurate and transparent.

47. *Pharmaceutical community relations* – This is the process of building relationships with the community. It involves participating in local events, supporting charitable causes, and addressing community concerns.

48. *Pharmaceutical environmental relations* – This is the process of managing the company's relationship with the environment. It involves reducing the company's carbon footprint, conserving resources, and addressing environmental concerns.

49. *Pharmaceutical social relations* – This is the process of managing the company's relationship with society. It involves promoting social responsibility, supporting social causes, and addressing social issues.

50. *Pharmaceutical corporate governance* – This is the process of overseeing the company's operations and ensuring that it is managed in a responsible and transparent manner. It involves setting standards, monitoring performance, and taking action when necessary.

51. *Pharmaceutical corporate social responsibility* – This is the process of ensuring that a company operates in a socially responsible manner. It involves addressing issues such as human rights, labor practices, and environmental impact.

52. *Pharmaceutical corporate citizenship* – This is the process of ensuring that a company is a good citizen of the community. It involves participating in local events, supporting charitable causes, and addressing community concerns.

53. *Pharmaceutical corporate social performance* – This is the process of measuring a company's social performance. It involves tracking and reporting on a range of social issues, such as human rights, labor practices, and environmental impact.

54. *Pharmaceutical corporate social impact* – This is the process of assessing the impact of a company's activities on society. It involves identifying the company's social responsibilities and measuring the impact of its actions.

55. *Pharmaceutical corporate social responsibility reporting* – This is the process of publishing a report that details a company's social responsibilities and performance. It is a key tool for communicating with stakeholders and demonstrating the company's commitment to social responsibility.

56. *Pharmaceutical corporate social responsibility strategy* – This is the process of developing a plan for how a company will address its social responsibilities. It involves identifying the company's social responsibilities and determining the actions that will be taken to address them.

57. *Pharmaceutical corporate social responsibility framework* – This is the process of establishing a set of principles and standards that guide a company's social responsibilities. It is a key tool for ensuring consistency and transparency in the company's social responsibilities.

58. *Pharmaceutical corporate social responsibility policy* – This is the process of creating a document that outlines a company's social responsibilities. It is a key tool for communicating the company's commitment to social responsibility.

59. *Pharmaceutical corporate social responsibility training* – This is the process of providing education and training to employees about the company's social responsibilities. It is a key tool for ensuring that all employees understand the company's commitment to social responsibility.

60. *Pharmaceutical corporate social responsibility audit* – This is the process of conducting an independent review of a company's social responsibilities. It is a key tool for ensuring the integrity of the company's social responsibilities reporting.

61. *Pharmaceutical corporate social responsibility certification* – This is the process of obtaining a certification from a third-party organization that recognizes a company's commitment to social responsibility. It is a key tool for demonstrating the company's commitment to social responsibility.

62. *Pharmaceutical corporate social responsibility award* – This is the process of receiving an award from a third-party organization that recognizes a company's commitment to social responsibility. It is a key tool for demonstrating the company's commitment to social responsibility.

63. *Pharmaceutical corporate social responsibility benchmarking* – This is the process of comparing a company's social performance to that of its peers. It is a key tool for identifying areas for improvement and setting goals for the future.

64. *Pharmaceutical corporate social responsibility benchmarking tool* – This is the process of using a tool to compare a company's social performance to that of its peers. It is a key tool for identifying areas for improvement and setting goals for the future.

65. *Pharmaceutical corporate social responsibility benchmarking data* – This is the process of collecting data on a company's social performance and comparing it to that of its peers. It is a key tool for identifying areas for improvement and setting goals for the future.

66. *Pharmaceutical corporate social responsibility benchmarking report* – This is the process of publishing a report that details a company's social performance and compares it to that of its peers. It is a key tool for identifying areas for improvement and setting goals for the future.

67. *Pharmaceutical corporate social responsibility benchmarking tool* – This is the process of using a tool to compare a company's social performance to that of its peers. It is a key tool for identifying areas for improvement and setting goals for the future.

68. *Pharmaceutical corporate social responsibility benchmarking data* – This is the process of collecting data on a company's social performance and comparing it to that of its peers. It is a key tool for identifying areas for improvement and setting goals for the future.

69. *Pharmaceutical corporate social responsibility benchmarking report* – This is the process of publishing a report that details a company's social performance and compares it to that of its peers. It is a key

Figure 1. The effect of the concentration of the *Agrobacterium* suspension on the transformation efficiency of *Agrobacterium* strains. The *Agrobacterium* strains were grown in the YEA medium for 24 h and then adjusted to the OD₆₀₀ of 0.1. The *Agrobacterium* strains were then grown in the YEA medium with the concentration of 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9, 1.0, 1.1, 1.2, 1.3, 1.4, 1.5, 1.6, 1.7, 1.8, 1.9, 2.0, 2.1, 2.2, 2.3, 2.4, 2.5, 2.6, 2.7, 2.8, 2.9, 3.0, 3.1, 3.2, 3.3, 3.4, 3.5, 3.6, 3.7, 3.8, 3.9, 4.0, 4.1, 4.2, 4.3, 4.4, 4.5, 4.6, 4.7, 4.8, 4.9, 5.0, 5.1, 5.2, 5.3, 5.4, 5.5, 5.6, 5.7, 5.8, 5.9, 6.0, 6.1, 6.2, 6.3, 6.4, 6.5, 6.6, 6.7, 6.8, 6.9, 7.0, 7.1, 7.2, 7.3, 7.4, 7.5, 7.6, 7.7, 7.8, 7.9, 8.0, 8.1, 8.2, 8.3, 8.4, 8.5, 8.6, 8.7, 8.8, 8.9, 9.0, 9.1, 9.2, 9.3, 9.4, 9.5, 9.6, 9.7, 9.8, 9.9, 10.0, 10.1, 10.2, 10.3, 10.4, 10.5, 10.6, 10.7, 10.8, 10.9, 11.0, 11.1, 11.2, 11.3, 11.4, 11.5, 11.6, 11.7, 11.8, 11.9, 12.0, 12.1, 12.2, 12.3, 12.4, 12.5, 12.6, 12.7, 12.8, 12.9, 13.0, 13.1, 13.2, 13.3, 13.4, 13.5, 13.6, 13.7, 13.8, 13.9, 14.0, 14.1, 14.2, 14.3, 14.4, 14.5, 14.6, 14.7, 14.8, 14.9, 15.0, 15.1, 15.2, 15.3, 15.4, 15.5, 15.6, 15.7, 15.8, 15.9, 16.0, 16.1, 16.2, 16.3, 16.4, 16.5, 16.6, 16.7, 16.8, 16.9, 17.0, 17.1, 17.2, 17.3, 17.4, 17.5, 17.6, 17.7, 17.8, 17.9, 18.0, 18.1, 18.2, 18.3, 18.4, 18.5, 18.6, 18.7, 18.8, 18.9, 19.0, 19.1, 19.2, 19.3, 19.4, 19.5, 19.6, 19.7, 19.8, 19.9, 20.0, 20.1, 20.2, 20.3, 20.4, 20.5, 20.6, 20.7, 20.8, 20.9, 21.0, 21.1, 21.2, 21.3, 21.4, 21.5, 21.6, 21.7, 21.8, 21.9, 22.0, 22.1, 22.2, 22.3, 22.4, 22.5, 22.6, 22.7, 22.8, 22.9, 23.0, 23.1, 23.2, 23.3, 23.4, 23.5, 23.6, 23.7, 23.8, 23.9, 24.0, 24.1, 24.2, 24.3, 24.4, 24.5, 24.6, 24.7, 24.8, 24.9, 25.0, 25.1, 25.2, 25.3, 25.4, 25.5, 25.6, 25.7, 25.8, 25.9, 26.0, 26.1, 26.2, 26.3, 26.4, 26.5, 26.6, 26.7, 26.8, 26.9, 27.0, 27.1, 27.2, 27.3, 27.4, 27.5, 27.6, 27.7, 27.8, 27.9, 28.0, 28.1, 28.2, 28.3, 28.4, 28.5, 28.6, 28.7, 28.8, 28.9, 29.0, 29.1, 29.2, 29.3, 29.4, 29.5, 29.6, 29.7, 29.8, 29.9, 30.0, 30.1, 30.2, 30.3, 30.4, 30.5, 30.6, 30.7, 30.8, 30.9, 31.0, 31.1, 31.2, 31.3, 31.4, 31.5, 31.6, 31.7, 31.8, 31.9, 32.0, 32.1, 32.2, 32.3, 32.4, 32.5, 32.6, 32.7, 32.8, 32.9, 33.0, 33.1, 33.2, 33.3, 33.4, 33.5, 33.6, 33.7, 33.8, 33.9, 34.0, 34.1, 34.2, 34.3, 34.4, 34.5, 34.6, 34.7, 34.8, 34.9, 35.0, 35.1, 35.2, 35.3, 35.4, 35.5, 35.6, 35.7, 35.8, 35.9, 36.0, 36.1, 36.2, 36.3, 36.4, 36.5, 36.6, 36.7, 36.8, 36.9, 37.0, 37.1, 37.2, 37.3, 37.4, 37.5, 37.6, 37.7, 37.8, 37.9, 38.0, 38.1, 38.2, 38.3, 38.4, 38.5, 38.6, 38.7, 38.8, 38.9, 39.0, 39.1, 39.2, 39.3, 39.4, 39.5, 39.6, 39.7, 39.8, 39.9, 40.0, 40.1, 40.2, 40.3, 40.4, 40.5, 40.6, 40.7, 40.8, 40.9, 41.0, 41.1, 41.2, 41.3, 41.4, 41.5, 41.6, 41.7, 41.8, 41.9, 42.0, 42.1, 42.2, 42.3, 42.4, 42.5, 42.6, 42.7, 42.8, 42.9, 43.0, 43.1, 43.2, 43.3, 43.4, 43.5, 43.6, 43.7, 43.8, 43.9, 44.0, 44.1, 44.2, 44.3, 44.4, 44.5, 44.6, 44.7, 44.8, 44.9, 45.0, 45.1, 45.2, 45.3, 45.4, 45.5, 45.6, 45.7, 45.8, 45.9, 46.0, 46.1, 46.2, 46.3, 46.4, 46.5, 46.6, 46.7, 46.8, 46.9, 47.0, 47.1, 47.2, 47.3, 47.4, 47.5, 47.6, 47.7, 47.8, 47.9, 48.0, 48.1, 48.2, 48.3, 48.4, 48.5, 48.6, 48.7, 48.8, 48.9, 49.0, 49.1, 49.2, 49.3, 49.4, 49.5, 49.6, 49.7, 49.8, 49.9, 50.0, 50.1, 50.2, 50.3, 50.4, 50.5, 50.6, 50.7, 50.8, 50.9, 51.0, 51.1, 51.2, 51.3, 51.4, 51.5, 51.6, 51.7, 51.8, 51.9, 52.0, 52.1, 52.2, 52.3, 52.4, 52.5, 52.6, 52.7, 52.8, 52.9, 53.0, 53.1, 53.2, 53.3, 53.4, 53.5, 53.6, 53.7, 53.8, 53.9, 54.0, 54.1, 54.2, 54.3, 54.4, 54.5, 54.6, 54.7, 54.8, 54.9, 55.0, 55.1, 55.2, 55.3, 55.4, 55.5, 55.6, 55.7, 55.8, 55.9, 56.0, 56.1, 56.2, 56.3, 56.4, 56.5, 56.6, 56.7, 56.8, 56.9, 57.0, 57.1, 57.2, 57.3, 57.4, 57.5, 57.6, 57.7, 57.8, 57.9, 58.0, 58.1, 58.2, 58.3, 58.4, 58.5, 58.6, 58.7, 58.8, 58.9, 59.0, 59.1, 59.2, 59.3, 59.4, 59.5, 59.6, 59.7, 59.8, 59.9, 60.0, 60.1, 60.2, 60.3, 60.4, 60.5, 60.6, 60.7, 60.8, 60.9, 61.0, 61.1, 61.2, 61.3, 61.4, 61.5, 61.6, 61.7, 61.8, 61.9, 62.0, 62.1, 62.2, 62.3, 62.4, 62.5, 62.6, 62.7, 62.8, 62.9, 63.0, 63.1, 63.2, 63.3, 63.4, 63.5, 63.6, 63.7, 63.8, 63.9, 64.0, 64.1, 64.2, 64.3, 64.4, 64.5, 64.6, 64.7, 64.8, 64.9, 65.0, 65.1, 65.2, 65.3, 65.4, 65.5, 65.6, 65.7, 65.8, 65.9, 66.0, 66.1, 66.2, 66.3, 66.4, 66.5, 66.6, 66.7, 66.8, 66.9, 67.0, 67.1, 67.2, 67.3, 67.4, 67.5, 67.6, 67.7, 67.8, 67.9, 68.0, 68.1

Figure 1 illustrates the experimental setup. A subject is seated at a table, looking at a video screen. A camera is positioned above the screen to capture the subject's hand position. A light source is positioned to the left of the screen. A target is positioned on the screen. The subject's hand is positioned near the target. The diagram shows the spatial arrangement of the subject, camera, screen, light source, and target.

the 1990s, the number of people in the world who are undernourished has declined from 1.1 billion to 800 million. The number of people who are malnourished has declined from 1.5 billion to 1 billion. The number of people who are obese has increased from 100 million to 300 million. The number of people who are overweight has increased from 100 million to 300 million. The number of people who are obese and overweight has increased from 100 million to 300 million. The number of people who are obese and overweight has increased from 100 million to 300 million.

In addition, a specific Operations Head Instruction requires monthly running of the standby CACs and the PAC for four hours (if not previously run during the month) to check mechanical operations.

3. Fail-Safe Testing of Safety-Related Pneumatic Equipment

Fail-safe testing of virtually all of Cook Nuclear Plant's air-operated, safety-related pneumatic equipment is accomplished either as a result of the Inservice Testing/Inspection (ISI) Program or the Technical Specification (TS) surveillance testing program. The few exceptions are discussed under the Section C.5 below. With respect to testing frequency, ASME Section XI requires quarterly testing of a valve's safety function under the fail-safe condition. The Cook Nuclear Plant's containment isolation valves are, for example, tested under this quarterly frequency; however, Technical Specifications impose a monthly test frequency for most of the valves within Section XI.

The safety-related, air-operated valves and dampers at Cook Nuclear Plant fail to their actuated position when air is removed from (or lost from) the pneumatic actuators. Fail-safe testing is accomplished by de-energization of a control air solenoid associated with each particular component. When the solenoids are de-energized, component motion to the actuated position is checked and timed (timing only applies to valves in the IST program). Cook Nuclear Plant's ISI personnel review and trend valve timing data determine which valves are not properly operating. These valves then undergo testing at an increased frequency or are repaired, if required. Fail-safe testing is also completed following any maintenance activity that potentially impacts valve performance. This was discussed earlier under Maintenance Practices in Attachment 3.

The plant's discrepancy reporting and job order system address the preventive and corrective actions with respect to failures of other pneumatic equipment. Failure positions of equipment other than valves, (e.g., air-operated dampers) were verified separately for this Generic Letter under C.6 below.

4. Inservice Testing Program

This section provides a description of the Inservice Testing Program that applies to the concerns of Generic Letter 88-14. Specifically, the testing program and test results for the air-operated valves, dampers and accumulators are presented. In addition, exceptions to the performance of fail-safe testing on certain air-operated equipment are provided following this section.



a. Valve Testing (General)

The Inservice Test Program (ISTP) for valves, as submitted to the NRC for the second ten-year inspection interval (Reference AEP:NRC:0969H dated October 5, 1987), commenced on August 23, 1985, and July 1, 1986 for Units 1 and 2, respectively. This valve test program is in accordance with Subsection IWV of Section XI of the 1983 edition of the ASME Boiler and Pressure Vessel Code through summer 1983 addenda, except for specific relief requests identified in the valve summary sheets of the program. The program includes the testing of all valve types (e.g., relief, safety, check, butterfly, gate, globe, diaphragm, etc.), and provides for the different valve actuator types (self-actuated check or relief, motor operated, air operated, solenoid operated, etc.). A valve summary sheet is used to identify the unique valve number as well as the valve position during both normal and post-accident plant operation. The sheets also indicate the primary test required by the code, the actual test performed, and the Technical Specification operating mode.

The discussion that follows applies only to air-operated safety-related valves as addressed by GL 88-14.

b. Valve Testing: Specific Response to Generic Letter 88-14

In order to respond to GL 88-14, a list of the safety-related air-operated valves in the Cook Nuclear Plant (188 in Unit 1 and 190 in Unit 2) was developed by reviewing the appropriate Cook Nuclear Plant drawings and updated FSAR Section. These valves were segregated according to the following:

- 1) Valves necessary to mitigate the consequences of postulated design bases accidents.
- 2) Valves expected to change position in response to a postulated accident as a result of an engineered safety features (ESF) actuation and not merely serve as a pressure boundary (such as pressurizer or steam generator PORVs). Note that, with a few exceptions, the fail-safe (loss of air) position is the "actuated" position, and these are discussed in C.5 below.

The safety-related, air-operated valves at Cook Nuclear Plant fail to their actuated position when air is removed from (or lost from) the pneumatic actuators. Periodic fail-safe testing is accomplished by de-energization of a control air

solenoid associated with each particular component as discussed earlier in Section C.3 of this attachment. This type of approach confirms the fail-safe function upon loss of power. It can also be concluded that this method verifies the loss of control air safety function by isolating the control air inlet flow path to the component and bleeding off air in the component actuators. Cook Nuclear Plant's ISI personnel review and trend valve timing data and determine which valves are not operating as expected. These valves then undergo testing at an increased frequency or are repaired as necessary. In specific response to the Generic Letter, the Cook Nuclear Plant's ISI personnel also reviewed the existing practice for evaluating the test results, and the bases for terminating increased frequency of surveillance. The review took into account significant performance attributes such as (1) control air solenoid valve conditions, (2) actuator diaphragm conditions, (3) limit switch adjustments, (4) actuator stem/valve stem coupling security, and (5) test parameters (including speed). This review by ISI personnel concluded that operating experience for the control air system has not indicated a problem to date due to valves failing to move to the safe condition on loss of air. However, the Cook Nuclear Plant has noted that ASCO solenoid valves have failed on occasions and it is recognized that this is an industry problem (reference: NRC IN 88-43), attributable to the valve diaphragm hardening. The Cook Nuclear Plant has a replacement program for these valves, by which they are rebuilt to ensure the proper functioning of these valves.

c. Valve Test Results

Unit 1

With the exception of seal leak testing on 10 valves scheduled for testing during the refueling outage in 1989 and those identified later under "Exceptions to Fail-Safe Testing," all 188 Unit 1 safety-related, air-operated valves were tested last in November 1988 (176 valves) and January 1989 (remaining 12 valves). Additionally, the latest testing of the diesel generators and their supporting startup air systems was conducted in February 1989 (D/G AB on February 20, 1989, and D/C CB on February 9, 1989).

[illegible]

Unit 2

All 190 Unit 2 valves (with the exception of those discussed under "Exceptions to Fail-Safe Testing") were last tested in March 1988 (50 valves), April 1988 (2 valves) and during the Steam Generator Replacement outage (138 valves). Additionally, the latest testing of the diesel generators (D/Gs AB and CD) and supporting startup air systems was conducted on February 13 and 20, 1989, respectively.

5. Exceptions to Fail-Safe Testing of Safety-Related Equipment and Components

The safety-related equipment and valves identified below are not fail-safe tested. The reasons and justifications for these exceptions are discussed below:

Virtually all of the safety-related, air-operated equipment fails to a safe position via solenoid action. The only exceptions are valves WRV-721 through -728; 1 and 2 XRV-221, -222, and -226 and -227; 1 and 2 XRV-220 and -225; and 1 and 2 CRV 470. The three-way regulating valves (WRV-721 through WRV-728) are located in the essential service water supply lines to the emergency diesel generators air aftercoolers. They regulate water flow to maintain the temperature at which the aftercooler air discharge thermostatic controller has been set. These valves are normally open and fail-safe open. They remain in service following an ESF actuation to regulate water flow. The valves cannot be stroke timed because they are thermostatic valves whose position is controlled by process fluid temperature and there is no external control available. A code relief request note on the testing requirements was submitted to the NRC under the inservice testing (IST) program for valves as an attachment to our letter to the NRC, AEP:NRC:0969, dated October 5, 1987.

It was discussed during this review for GL 88-14, that the valves WRV-721 through -728 have not been fail-safe tested as stated in the IST program for valves. Appropriate corrective action is being taken to evaluate and rectify this deficiency.

The reasons for the code relief request are that (1) these valves function only as regulating valves and not open/closed valves, (2) these valves are demonstrated operable during diesel generator testing (diesel generators are tested on a staggered basis every 31 days per Technical Specification 4.8.1.1.2), and (3) these valves are demonstrated operable during diesel generator 24-hour runs performed each refueling outage. The valves will be "fail-safe" tested during refueling outages. These valves on Unit 2 (WRV-722, 724, 726 and 728) were last tested for proper functioning in

1. The first part of the report is a general introduction to the subject of the study. It discusses the importance of the study and the objectives of the research.

2. The second part of the report is a detailed description of the methodology used in the study. It includes information about the sample size, the data collection methods, and the statistical analysis techniques.

3. The third part of the report is a presentation of the results of the study. It includes a summary of the findings and a discussion of the implications of the results.

4. The fourth part of the report is a conclusion and a list of references. The conclusion summarizes the main findings of the study and provides a final statement on the research.

5. The fifth part of the report is a list of references. It includes a list of all the sources used in the study, including books, articles, and other documents.

6. The sixth part of the report is a list of appendices. It includes a list of all the additional materials that are included in the report, such as tables, figures, and other documents.

7. The seventh part of the report is a list of footnotes. It includes a list of all the footnotes that are included in the report, providing additional information on the research.

8. The eighth part of the report is a list of acknowledgments. It includes a list of all the people and organizations that have provided support and assistance during the course of the study.

January 1989 during the refueling outage. The valves on Unit 1 (WRV-721, -723, -725 and -728) were last tested for proper functioning in January 1989 during the diesel generators' monthly surveillance running.

Additionally, the following air-operated valves are tested, as required by the code, under the Inservice Testing Program for valves, for operability (i.e., exercised full stroke) every three months but are not fail-safe tested for the reasons discussed below. Also, a code relief request for not performing the required stroke time testing of 10 seconds was made under the Cook Nuclear Plant's Inservice Testing Program for valves submitted to the NRC in October 1987 (letter to the NRC, AEP:NRC:0969).

The EDG turbo assist valves, XRV-220 on engine AB of each unit and XRV-225 on engine CD of each unit are normally closed and fail closed. They are required to open when the EDGs start since turbo jet assist is necessary to satisfy the fast start (10 seconds) capability of the EDGs. Testing of the turbo assist valves to verify their safety-related actuation is, therefore, conducted when the EDGs are Technical Specification surveillance tested on a 30-day staggered test basis. Given that the safety function of the turbo assist valves is to open, no fail-safe testing is necessary for these valves. In addition to the code required operability test stated above, these valves are also exercised (with fail-safe actuators) every three months in compliance with the code. These code tests were last performed on a staggered basis on both valves of both engines in January 1989.

The EDG starting air supply valves, XRV-221 and XRV-222 on engine AB of each unit and XRV-226 and XRV-227 on engine CD of each unit. These normally closed valves fail "as-is" and are demonstrated operable on every other EDG Technical Specification surveillance test, (see Section 4.8.1.1.2). The testing for each valve is conducted during the EDG start by isolating the accumulator associated with each of these valves on an alternating basis once per quarter. If these valves were to fail in other than the closed direction, the EDG would start. For these reasons, and since these valves fail "as-is", no fail-safe testing is performed. The code required operability testing of these valves every three months, as stated above, was last performed as follows:

- o Unit 1, AB diesel engine in November 1988, and CD engine in January 1989.
- o Unit 2, AB diesel engine in January 1989, and CD engine in November 1988.

1. The first part of the report discusses the importance of maintaining accurate records of all transactions. It emphasizes that this is crucial for the company's financial health and for providing reliable information to stakeholders. The report also mentions that the records should be kept up-to-date and that any discrepancies should be identified and corrected immediately.

2. The second part of the report focuses on the need for transparency in the company's operations. It states that transparency is essential for building trust with customers and investors. The report suggests that the company should provide regular updates on its performance and that it should be open to external audits to ensure that its financial statements are accurate and reliable.

3. The third part of the report discusses the importance of maintaining a strong relationship with the company's suppliers. It notes that a good relationship with suppliers can help the company to secure better prices and to ensure that its supply chain is stable. The report also mentions that the company should communicate regularly with its suppliers and that it should be willing to negotiate terms that are fair and reasonable for both parties.

4. The fourth part of the report discusses the importance of maintaining a strong relationship with the company's customers. It notes that a good relationship with customers can help the company to increase its sales and to ensure that its products are of high quality. The report also mentions that the company should communicate regularly with its customers and that it should be willing to provide excellent customer service.

5. The fifth part of the report discusses the importance of maintaining a strong relationship with the company's employees. It notes that a good relationship with employees can help the company to attract and retain top talent. The report also mentions that the company should communicate regularly with its employees and that it should be willing to provide a safe and healthy work environment.

6. The sixth part of the report discusses the importance of maintaining a strong relationship with the company's shareholders. It notes that a good relationship with shareholders can help the company to raise capital and to ensure that its stock price is high. The report also mentions that the company should communicate regularly with its shareholders and that it should be willing to provide accurate and reliable financial information.

Component cooling water valves 1 and 2 CRV-470. A code relief request (AEP:NRC:0969H) has been made under the IST valve program for air-operated valves 1- and 2-CRV-470, which are located in the component cooling water (CCW) return from the letdown heat exchanger of each unit. These valves control the temperature of the letdown flow leaving the heat exchanger. These valves cannot be "fail safe" tested nor stroke timed since no control switches are installed to perform those tests. These valves are full stroke exercised quarterly. These valves were last full stroke exercised in January 1989 and December 1988 on Units 1 and 2, respectively.

An additional piece of equipment for which fail-safe testing is not performed is the governor for the turbine driven auxiliary feedwater pump (TDAFP). Although fail-safe testing on this equipment is not performed, the TDAFPs are exercised monthly in Modes 1, 2, and 3 using the associated unit's TDAFP test procedures. In addition, any controllability problems with the governor could be overcome by using the control room tripping capability and/or local manual control. Motor-operated valves (MOVs) in the discharge of this pump would also enable the operators to throttle the TDAFP flow to the steam generators. All these factors were evaluated as a result of the NRC's IE Inspection Reports 50-315/86-13 and 50-316/84-15, and fail-safe testing of this component was not considered warranted.

6. Safety-Related, Air-Operated Dampers Testing

There are 12 safety-related, air-operated dampers which are surveillance tested on a monthly basis in accordance with a Plant Operation Head Technical Specification Surveillance Test Procedure. These include six dampers (3 for each train - 1-HVAES-1D1 through 1D3 and 1-HVAES-2D1 through 2D3) for Unit 1 in the auxiliary building ESF ventilation system. Six other dampers serve a similar purpose for Unit 2. The first two in each train are normally open and bypass the charcoal filters, and fail-safe to close. The third damper (i.e., 1-HVAES-1D3 and 1-HVAES-2D3), which is normally closed, is on the face of the charcoal filter on each train and it fails safe open. The dates that these dampers were last tested are January 15 and January 28, 1989, for Units 1 and 2, respectively.

7. Pneumatic Accumulators Design Verification

In response to NRC IN 87-28, a review was conducted of air-operated valves equipped with air accumulator bottles. The review took into account pre-operational test procedures for loss of instrument air (both units), the plant's previous safety-related design changes, system descriptions, flow

THE
FEDERAL
BUREAU OF
INVESTIGATION
UNITED STATES DEPARTMENT OF JUSTICE
WASHINGTON, D. C. 20535

MEMORANDUM FOR THE DIRECTOR

SUBJECT: [Illegible]

[Illegible text follows, appearing to be a memorandum format with a subject line and several paragraphs of text that are mostly illegible due to the quality of the scan.]

[Illegible text continues, appearing to be the body of a memorandum or report, with several paragraphs of text that are mostly illegible due to the quality of the scan.]

diagrams, physical drawings and AEPSC's response to NRC IN 85-35 ("Failure of Air Check Valves to Seat"). Both safety-related and non safety-related accumulators were identified during the review.

The system valves identified as having permanently installed, non safety-related air accumulators were: condenser steam dump valves (URVs); feedwater regulating valves (FRVs); non essential service water (NESW) supply and return containment isolation valves (WCRs); ice condenser glycol supply and return containment isolation valves (VCRs); and containment ventilation supply and exhaust containment isolation valves (VCRs). Upon loss of control air, the above valves go to their fail-safe position which is closed. The URVs are non safety-related. Since the safety function of the FRVs, WCRs, and VCRs is to close, the air accumulators do not provide back-up to perform that safety function. The concerns of Recommendation #3 to Generic Letter 88-14, therefore, do not apply to these valves.

The review also included verification of the adequacy of safety-related air-operated valves for the functioning of pressurizer power operated relief valves PORVs while in Mode 5 for low temperature/overpressurization protection (LTOP) of the reactor vessel and the RCS. One set of safety-related air-operated valves, NRV-152 and -153 (pressurizer power-operated relief valves (PORVs)) are equipped with backup air accumulators as well as with air storage receivers. The operability of the PORV accumulators is verified periodically in accordance with Technical Specification requirements.

Air storage receivers were also identified on the emergency diesel generator starting air system, which is described in Attachment 1. The starting air system is tested periodically in conjunction with emergency diesel generator testing required by the Technical Specifications.

The review conducted for NRC IN 87-28 concluded that the non safety-related air receivers provided in the plant are adequate for operational consideration. Further, the LTOP redundant air supply is considered adequate to assist the PORVs to open within the required LTOP stroke time and remain functional during or after an OBE (operational basis earthquake). Finally, the emergency diesel generator starting air system receivers are of an adequate design as verified by pre-operational testing and functionality of the system is demonstrated by the monthly surveillance testing.

D. Verification of Correct Failure Positions

The inservice testing program for valves contains both failure and safety positions of all air-operated, safety-related valves in flow paths associated with engineered safety feature systems. These valve positions were verified correct during development of the program.

