10 CFR 50.73

PHILADELPHIA ELECTRIC COMPANY

LIMERICK GENERATING STATION

P. O. BOX A

SANATOGA, PENNSYLVANIA 19464

(215) 327-1200 EXT. 2000

June 22, 1989

M. J. MCCORMICK, JR., P.E. PLANT MANAGER LIMERICK GENERATING STATION

Docket No. 50-352 License No. NPF-39

U.S. Nuclear Regulatory Commission Attn: Document Control Desk Washington, DC 20555

> SUBJECT: Licensee Event Report Limerick Generating Station - Unit 1

This LER reports a condition prohibited by Technical Specifications in that pressure, flow, and level indication was incperable due to environmentally ungualified valve seals.

Reference:	Docket No. 50-352
Report Number:	89-034
Revision Number:	00
Event Date:	May 12, 1989
Discovery Date:	May 22, 1989
Report Date:	June 22, 1989
Facility:	Limerick Generating Station
	P.O. Box A, Sanatoga, PA 19464

This LER is being submitted pursuant to the requirements of 10 CFR 50.73(a)(2)(i)(B) and 50.73(a)(2)(v)(D). This LER is being submitted one day late because of delays in administrative processing. We regret any inconvenience this may have caused.

Very truly yours, m. m. Cosmit

WGS:sc

cc: W. T. Russell, Administrator, Region I, USNRC T. J. Kenny, USNRC Senior Resident Inspector, LGS

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accordance with 10CFR50.72(b)(2)(iii)(D) due to a condition that could have prevented the fulfillment of a safety function of systems that are needed to mitigate the consequences of an accident.

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Plant Conditions Prior to the Event:

Operating Mode: 5 (Refueling)

Reactor Power: 0%

Description of the Event

The Limerick Generating Station Final Safety Analysis Report (FSAR), Chapter 1, "Introduction of General Description of the Plant," Section 1.8, states that the requirements of Regulatory Guide (R.G.) 1.97, Revision 2, 1980, "Instrumentation for Light-Water-Cooled Nuclear Power Plants to Assess Plant Conditions During and Following an Accident," are met to the extent discussed in FSAR Section 7.5. R.G. 1.97 identifies the plant variables to be measured and the instrumentation criteria for assuring acceptable emergency response capabilities during and following a Design Basis Accident (DBA).

During June 1988, as the result of a review of surveillance test methods on Unit 2, a request by station personnel for the addition of test valves (EIIS:TV) in Unit 2 instrument lines (similar to Unit 1) was initiated. In response to this request, in December 1988, it was found by Engineering that the one-hundred nine (109) NUPRO brand valves that had been installed on Unit 1 instrument lines during the Unit 1 preoperational testing phase prior to receipt of the Low Power Operating License were non-ASME code valves. Ninety-nine (99) were installed on General Electric (GE) supplied local instrument racks outside the ASME code boundaries and ten (10) were installed on non-GE supplied level instrument racks within the ASME code boundaries. The test valve installations in question were designed to allow simulation of a lie break in the instrument lines in order to functionally check the primary containment excess flow check valves.

In January 1989, Philadelphia Electric Company (PECo) requested that the Architect Engineer (A/E) initiate a Modification Design Change Package (MDCP) to revise ASME code boundaries and pipe classifications for Unit 1 that would have allowed the installation of the subject test valves. An Engineering Work Request (EWR) was initiated in March 1989 to obtain formal engineering resolution of the issue. At the end of April 1989 a Nonconformance Report (NCR) was issued to formally identify the

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non-ASME test values on ASME code lines as an item to be resolved. Subsequent investigation then revealed that the Unit 1 test values had not been procured in accordance with design specifications. On May 6, 1989 a second NCR was issued and formally identified that the test values were not Qualified (Q) but were installed in Q-passive applications (i.e., Q-listed item however its only function is to maintain the required pressure boundary).

On May 12, 1989, the plant staff was notified by the A/E that the ninety-nine (99) test valves installed on non-GE supplied instrument racks satisfied the ASME code boundary criteria as defined in ASME Section XI, IWA 7000. For the ninety-nine (99) NUPRO valves installed on General Electric rack mounted instruments, the valves were originally installed beyond the ASME boundaries and therefore the ASME code does not apply to these valves. However, an environmental qualification problem was identified with the soft parts of the NUPRO valves. The test valves contained teflon coated viton seals (EIIS:SEAL) as part of the valves pressure boundary. The A/E calculated that the radiation dose that would be received by the test valves eight (8) hours after a DBA Loss of Coolant Accident (LOCA) would cause degradation of the valve seals leading to the potential to allow leakage causing inaccurate instrument indications. As a result of the potential for the loss of the required instrumentation due to breakdown of the teflon coated viton seals, nineteen (19) test valve installations were removed on May 13, 1989. The test valves had been installed on various instrument lines and fifteen (15) of nineteen (19) test valves were installed on instrumentation for vessel level (EIIS:LI) and pressure indications (EIIS: PI), Main Steam Line Isolation Valve (MSIV) Leakage Control (EIIS: BD) system pressure indications, and reactor recirculation pump (EIIS:P) flow indications (EIIS:FI). These indications are required to be operable longer than eight (8) hours following a DBA LOCA to comply with R.G. 1.97. The purpose of this instrumentation is to assure acceptable emergency response capabilities during and following the course of an accident. In addition, four (4) test valves associated with the Reactor Core Isolation (RCIC)(EIIS:BN) system steam supply instrumentation were also replaced as a conservative measure. This instrumentation is only required to be operable for one (1) hour following the DBA LOCA and the instrumentation would be unaffected by the condition of the test valve seals.

On May 22, 1989, plant staff identified that six (6) of the nineteen test valves were located on instrument lines that are connected to instrumentation required by R.G. 1.97 and Technical

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Specifications (TS). The unqualified test valves affected the operability of these six (6) instruments and TS section 3.3.7.5 requires that a minimum number of one channel of reactor vessel pressure and level instrumentation be operable. However, this TS was not met due to the environmentally unqualified test valve seals.

This condition has existed since October 26, 1984, the date of the issuance of the Unit 1 Low Power Operating License. The "Action" required by TS Section 3.3.7.5 was not taken in the specified time period, therefore, for the six (6) test valves associated with the TS required instrumentation this constitutes a condition prohibited by TS and is reportable in accordance with 10CFR 50.73(a)(2)(i)(B). Additionally, for the fifteen (15) test valves associated with the instrumentation required by R.G. 1.97 this constitutes a condition that could have prevented the fulfillment of a safety function of systems that are needed to mitigate the consequences of an accident and is reportable in accordance with 10CFR50.73(a)(2)(v)(D).

In addition, upon further evaluation, plant staff determined that this event resulted in a condition that could have prevented the fulfillment of the safety function of systems and components that are needed to mitigate the consequences of an accident, and a four-hour notification to the NRC was made on May 24, 1989 at 1615 hours in accordance with the requirements of 10CFR 50.72(b)(2)(iii)(D).

Consequences of the Event:

The actual consequences of this condition were minimal because a DBA LOCA that could result in degradation of the test valve pressure boundary seal did not occur and the installation of these valves does not impact normal plant operations. There was no release of radioactive material to the environment as a result of this condition.

The potential consequences of this condition were that instrumentation for Reactor level and pressure, MSIV Leakage Control system pressure and Reactor Recirculation pump flow may not have provided the accurate indications necessary to assess equipment and plant conditions following a DBA LOCA. The Control Room operator would utilize the post accident qualified instrument indications to monitor transient reactor plant behavior and to verify proper safety system performance following

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the accident. Approximately eight (8) hours following the DBA LOCA, the test valve seals may start to leak due to radiation exposure degradation, therefore, making post accident qualified instruments inaccurate. The inaccurate indications could affect the ability of the Control Room operators to verify adequate reactor coolant inventory and pressure, adequate MSIV Leakage Control system operation and Reactor Recirculation pump flow.

The leaking test valve seals would have also resulted in contamination of certain areas of the Reactor Enclosure (EIIS:NG). However, personnel access to the Reactor Enclosure would be restricted after the DBA LOCA due to expected radiation levels resulting from the accident regardless of the condition of the test valve seals. The contaminated fluid that leaked from the valves would be processed by the floor drain system and the Reactor Enclosure Recirculation system (EIIS:VA) (RERS) and Standby Gas Treatment system (EIIS:BH) (SGTS) would process the airborne contamination.

In summary, if a DBA LOCA had occurred, the radiation induced degradation of the NUPRO valve seals and resultant leakage could have adversely affected the ability of the plant and Control Room operators to mitigate the consequences of the accident.

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Cause of the Event:

This event was a result of an inadequate review of those system design specifications against the materials (i.e. valves) intended for installation on the part of plant staff supervision. The valves located on GE supplied local instrument racks were installed during the Unit 1 preoperational testing phase prior to receipt of the Low Power Operating License with the approval of General Electric, the Nuclear Steam Supply System (NSSS) supplier, under Field Deviation Disposition Request (FDDR) HH1-4167. The FDDR approved valve installation but left material qualification and procurement of proper materials (i.e. valves) as the Philadelphia Electric Company's responsibility. This responsibility was not recognized. In addition, the valves installed on non-GE supplied local instrument racks were also installed prior to receipt of the Low Power Operating License using the guidance supplied by the FDDR, but without formal design approval by the A/E due to a similar error on the part of the plant staff supervision.

In summary, the installation of these valves failed to comply with either A/E or the PECo Engineering Department modification procedures in effect at the time due in part to the multiple modification procedures in effect prior to receipt of the low power license.

Corrective Actions:

All nineteen test valves required to be removed were removed by Instrumentation and Control technicians under Maintenance Request Forms by May 13, 1989. In response to the disposition to the Nonconformance Reports the additional one-hundred and one valves left in place will be administratively controlled as follows:

- A routine test, RT-2-000-631-1, was written to perform required leak inspections of remaining NUPRO instrument test valves until the third Unit 1 refuel outage.
- The number of valve cycles is being tracked through Instrument and Control Department aid tags hung on each remaining NUPRO test valve.
- 3) Plant staff is initiating Modification requests to remove and replace the remaining valves prior to restart from the Unit 1 third refueling outage.

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Actions Taken to Prevent Recurrence:

Upon receipt of the Unit 1 Low Power Operating License, Administrative Procedure A-14 "Procedure for Control of Plant Modifications", was placed in effect and adequately provided instruction and control throughout the modification process. This procedure addresses the modification review process and involves the independent review of several specialized work groups, supervision and management. Plant staff has determined that the current modification process is adequate and provides the proper instruction to attain the appropriate independent reviews. A detailed root cause analysis will be performed by August 31, 1989, to determine whether further investigation into the generic concern is necessary.

Previous Similar Occurrences:

None

Cause	Code:	D2	Inadequate	Procedure	
		A2	Failure to	follow implementing p	rocedures

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