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TELEPHONE ARCA CODE 710 546-2700

February 3, 1992

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

Subject:

LER 92-001, Failure of Containment Radiation Monitor Due To Unknown Cause, Causes Containment Ventilation

Isolation (i.e. ESF Actuation) R.E. Ginna Nuclear Power Plant

Docket No. 50-244

In accordance with 10CFR50.73, Licensee Event Report System, item (a)(2)(iv), which requires a report of, "any event or condition that resulted in manual or automatic actuation of any Engineered Safety Feature (ESF), including the Reactor Protection System (RPS)", the attached Event Report LER 92-001 is hereby submitted.

This event has in no way affected the public's health and safety.

Very truly yours,

Robert C. Mecredy

xc:

U.S. Nuclear Regulatory Commission

Region I

475 Allendale Road

King of Prussia, PA 19406

Ginna USNRC Senior Resident Inspector

Cet No 1845931141

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On January 5, 1992 at approximately 0240 EST, with the reactor at approximately 98% full power, a containment ventilation isolation occurred due to an actuation signal from the containment particulate radiation monitor (R-11).

All containment isolation valves that were open, closed as designed.

Immediate operator action was to perform the applicable alarm response procedures actions. This included verifying automatic actions, determining the cause of the containment ventilation isolation, and making appropriate notifications.

The immediate cause of the event was determined to be the failure of R-11.

Corrective action taken was to return the containment ventilation isolation system to pre-event normal status, sequentially followed by a troubleshooting effort by the Instrument and Control Department, and then changeout of the R-11 drawer with a qualified spare. Further investigation to determine the root cause is continuing.

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I. PRE-EVENT PLANT CONDITIONS

The plant was at approximately 98% steady state reactor power with no major activities in progress.

II. DESCRIPTION OF EVENT

DATES AND APPROXIMATE TIMES OF MAJOR OCCURRENCES:

- January 5, 1992, 0240 EST: Event date and time.
- January 5, 1992, 0240 EST: Discovery date and 0 time.
- January 5, 1992, 0252 EST: Control Room operators restore R-11 (Containment Particulate Radiation Monitor and reset containment ventilation isolation).

B. EVENT:

On January 5, 1992 at approximately 0240 EST, with the reactor at approximately 98% full power, following control board alarms were received, E-16 (RMS Process Monitor High Activity) and A-25 (Containment Ventilation Isolation). The Control Room operators, responding to the above alarms, observed that R-11 (Containment Particulate Radiation Monitor) had the light indicating failure illuminated. Control Room operators immediately referred to alarm response procedures AR-A-25 and AR-RMS, and verified that all containment ventilation isolation valves that were open, closed as designed and performed the applicable actions of the alarm response procedures. Subsequently, at approximately 0242 EST, Control Board alarm E-20 (CNMT Or Plant Vent Rad Mon Pump Trip) was received. This alarm was due to the trip of the containment radiation monitor pump and isolation of the containment valves to and from the pump. The Control Room operators also verified that the other containment process radiation monitors were reading normal prior to the radiation monitor pump trip.

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After the above immediate actions were completed, the Control Room operators addressed plant Technical Specifications and declared R-11 inoperable.

At approximately 0252 EST, January 5, 1992, the Control Room operators reset R-11 by cycling its AC power supply off and on, reset the containment ventilation isolation signal, and restarted the containment radiation monitor pump. All containment process radiation monitor readings returned to approximately pre-event values, indicating that R-11 was now operating properly. Subsequently, at 0324 EST, the Control Room operators performed periodic test procedure PT-17.2 (Process Radiation Monitors R-11 - R-22 Iodine Monitors R-10A and R-10B) on R-11 only and demonstrated that R-11 was operating as required.

C. INOPERABLE STRUCTURES, COMPONENTS, OR SYSTEMS THAT CONTRIBUTED TO THE EVENT:

None.

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D. OTHER SYSTEMS OR SECONDARY FUNCTIONS AFFECTED:

With the containment ventilation isolation, the following major components were isolated:

- o R-10A, Containment Iodine RMS Monitor
- o R-11, Containment Particulate RMS Monitor
- o R-12, Containment Gas RMS Monitor

E. METHOD OF DISCOVERY:

The event was immediately apparent due to Control Board annunciator alarms and containment ventilation isolation valve position indication on the Control Board. Also, Radiation Monitor R-11 digital readout indicated an invalid error code.

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F. OPERATOR ACTION:

Control Room operators responded to the event by performing the applicable actions of alarm response procedures E-16, A-25, RMS, and E-20 and other actions as they deemed necessary. This included the following:

- Verifying that all containment ventilation isolation valves that were open, closed as designed.
- Addressing the plant Technical Specifications to ensure the plant was operating within these specifications.
- o Declaring R-11 inoperable per administrative procedure A-52.4 (Control of Limiting Conditions for Operating Equipment).
- o Resetting R-11, resetting the containment ventilation isolation signal and restarting R-10A, R-11, and R-12 sample pump and verifying sample flow was re-established.
- O Verifying that R-10A, R-11, R-12 RMS monitor readings returned to normal.
- o Notifying the NRC and higher supervision of the ESF actuation.

G. SAFETY SYSTEM RESPONSES:

The containment ventilation isolation valves that were open, closed automatically from the containment ventilation isolation signal.

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III. CAUSE OF EVENT

A. IMMEDIATE CAUSE:

The containment ventilation isolation was due to an R-11 failure.

B. ROOT CAUSE:

After the following troubleshooting, the root cause still remains undetermined at this time:

- The Instrument and Control (I&C) Department calibrated the R-11 drawer with no adjustments required.
- Victoreen Inc., the manufacturer of the instrument was called. Victoreen Inc. concluded that, the probable cause was the micro-processor "lockingup" and it was reset by the operators cycling its AC power supply off and on. They suspect it may be a "one time" event.

IV. ANALYSIS OF EVENT

This event is reportable in accordance with 10CFR50.73, Licensee Event Report system, item (a)(2)(iv), which requires reporting of, "any event or condition that resulted in manual or automatic actuation of any Engineered Safety Feature (ESF) including the Reactor Protection System (RPS)". The containment ventilation isolation due to the R-11 failure, was an automatic actuation of an ESF subsystem.

An assessment was performed considering both the safety consequences and implications of this event with the following results and conclusions:

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There were no operational or safety consequences or implications attributed to the containment ventilation isolation because:

- The containment ventilation isolation system operated as designed.
- The components affected were capable of withstanding the isolation.
- The containment ventilation isolation was in the conservative direction.

Based on the above, it can be concluded that the public's health and safety was assured at all times.

V. CORRECTIVE ACTION

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- ACTION TAKEN TO RETURN AFFECTED SYSTEMS TO PRE-EVENT NORMAL STATUS:
 - The Control Room operators, after determining that the containment ventilation isolation was due to the R-11 failure, reset R-11, reset the containment ventilation isolation signal and restored the system to pre-event status.

ACTION TAKEN OR PLANNED TO PREVENT RECURRENCE: B.

The following corrective action was taken:

- The R-11 drawer was replaced with a qualified spare and the removed R-11 drawer will be sent to Victoreen, Inc., so that they can attempt to duplicate the failure and determine the root cause.
- Engineering has been involved in assessing the situation and will provide guidance for any desirable follow-up actions.

No other corrective action is planned until a root cause determination is accomplished.

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VI. ADDITIONAL INFORMATION

A. FAILED COMPONENTS:

The R-11 drawer was a model #942A, manufactured by Victoreen, Inc.

PREVIOUS LERS ON SIMILAR EVENTS: B.

A similar LER event historical search was conducted with the following results: LERs 87-005, 88-007, 89-011, 89-013, and 89-014 were similar events with known causes that appear much different than this event. No other documentation of similar events could be identified.

C. SPECIAL COMMENTS:

None.