
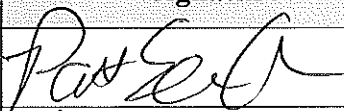

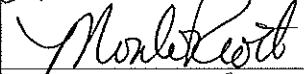
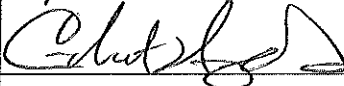
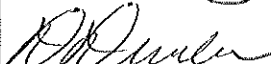
	GILES CHEMICAL ~ PREMIER MAGNESIA		
	Validation Protocol		
	Title: Scales IQ/OQ Protocol	Number: E13-VAL-RIQ-651	
	Owner: Patrick Owen	Revision: 0	
	Effective Date: June 19, 2013	Page: 1 of 12	

Approvals

Signing below indicates agreement that the protocol is ready for execution of the Installation and Operational Qualification for the Air Compressors located at 396 Smathers Street in Waynesville, NC.

Project Team Member	Functional Area	Signature	Date
Patrick Owen	Engineering		6/19/2013
Robert Willis <i>Lester Panton</i> per 6/19/2013	Maintenance		6-19-13
Monte Plott	Production		6-19-13
Matt Haynes	Operations		6-19-13
Deborah Durbin	Quality		6-19-13

A final summary report that consists of results and conclusions based on the data collected after protocol execution will be written and approved. The executed protocol will be attached behind the report.

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I. PURPOSE:

The purpose of this protocol is to certify with documented evidence that the Repackaging Scales are installed and function as intended. This protocol sets forth the objectives, methodology, documentation, and test activities needed to complete the Installation Qualification (IQ) and Operational Qualification (OQ) for the Scales located at Giles Chemical Repackaging Unit, 396 Smathers Street, Waynesville, NC.

II. BACKGROUND:

Giles' products are sold by weight. In order to verify that the machinery is packaging the correct amount of Epsom Salt into each package, scales are used to verify weights. The results are then used to adjust the machinery controls to dispense the correct amount of salt into the package. Giles has 4 scales in the Repackaging Area that are used for these process control purposes.

The products that are impacted by this study are all Epsom Salt products manufactured by Giles Chemical.

III. OVERVIEW

No other departments or systems will be affected by the installation or use of this equipment.

The following tests will be performed in this qualification:

Installation Documentation – the serial number or asset tag number of each scale will be documented.

Utility Verification – the voltage to each scale will be documented and verified to be correct.

Control / Operation Verification – the controls will be verified

IV. SYSTEM DESCRIPTION:

A. The system consists of 4 scales.

B. Description of Operation

01. The scales are started by pressing the power button.

02. The scales are tared by pressing the tare button.

03. The product is weighed.

04. For weight checks, an empty package is placed on the scale before taring. The weight measured of the filled package is then assumed to be the weight of the Epsom salt dispensed.

V. SCOPE



The Installation and Operational Qualification protocol is intended to certify with documented evidence that the scales are installed properly and function as desired by Giles..

VI. ROLES AND RESPONSIBILITIES

1. Engineering

- ❖ Write and issue the protocol
- ❖ Investigate protocol deviation reports

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- ❖ Execute the IQ and OQ.
 - ❖ Review raw data and originate interim notification to Quality Assurance
 - ❖ Write and route the final report
2. Quality Assurance
- ❖ Review and approve the protocol.
 - ❖ Review and approve raw data and notifications.
 - ❖ Review, approve, and store the final report.
3. Maintenance
- ❖ Provide Equipment Manuals, if available, to execute operational qualification.
 - ❖ Review and approve the protocol.
 - ❖ Assist with executing the IQ and OQ if needed.
 - ❖ Review and approve raw data and notifications.
 - ❖ Review and approve the final report
4. Production
- ❖ Review and approve the final report.

VII. TEST PROGRAM

A. INSTALLATION QUALIFICATION

Objective

The objective of the installation verification is to document each piece of Air Compressor equipment.

Equipment/Materials

Scales #1 - #4



Ideal Digital Multimeter Model #61-340 (SN 100100221)

Procedure

Perform each listed below for each scale

- Location: Verify that the equipment is situated to allow sufficient room around the machine for access doors and panels to be opened.
- Level – place a level on the scale and verify that the scale is level.
- Equipment: Document the Model and Serial or Asset Tag number of each piece of each scale
- Utilities
 - Electrical Requirements: Verify that instrument is receiving its specified Voltage.

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Acceptance Criteria

If the voltage is correct, each piece is uniquely identified, and sufficient access for all doors and panels is available, the scales will be considered installed properly.

B. OPERATION QUALIFICATION

Objective

The objective of Controls Verification is to document that the Scales operate as needed by Giles. The controls will be operated to test the ability of the Scales to be started and tared as needed.

Equipment/Materials

Scale #1 - #4

Procedure

Start each scale with the power button.

Place a weight on the scale and press the tare button. Verify that the scale tares properly.

Acceptance Criteria

If each scale powers up and tares properly the scales will be considered operationally qualified.

VIII. CALIBRATION

Verify that all instrumentation that requires calibration is calibrated.

- Ideal Digital Multimeter Model #61-340 (SN 100100221)
- Scale #1
- Scale #2
- Scale #3
- Scale #4

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**Scales: INSTALLATION QUALIFICATION****A. Installation Qualification****01. Location****a. Scale #1:**

LOCATION			
Distance Criterion	Does the Scale meet the criteria?(Yes/No)	Verified By	Date
Allow sufficient room around the machine for access doors and panels to be opened			
The scale should be level			

b. Scale #2:

LOCATION			
Distance Criterion	Does the Scale meet the criteria?(Yes/No)	Verified By	Date
Allow sufficient room around the machine for access doors and panels to be opened			
The scale should be level			

c. Scale #3:

LOCATION			
Distance Criterion	Does the Scale meet the criteria?(Yes/No)	Verified By	Date
Allow sufficient room around the machine for access doors and panels to be opened			
The scale should be level			

d. Scale #4:

LOCATION			
Distance Criterion	Does the Scale meet the criteria?(Yes/No)	Verified By	Date
Allow sufficient room around the machine for access doors and panels to be opened			
The scale should be level			

Reviewed By: _____

Date: _____

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**02. Equipment Identification**

Equipment Identification			
Equipment	Serial or Tag Identifier	Verified By	Date
Manual Line #1			
Scale #1			
Scale #2			
Scale #3			
Scale #4			
Comments:			

03. Utilities

- a. Verify that unit is receiving its specified utility requirements.



Electrical			
Specified	Actual	Verified By	Date
110-120 V Scale #1			
110-120 V Scale #2			
110-120 V Scale #3			
110-120 V Scale #4			
Comments:			

Reviewed By: _____

Date: _____

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Air Compressor: OPERATIONAL QUALIFICATION

B. Operation Qualification

01. Controls Verification – to document that the scale controls work properly



Controls/Indicators Verification				
Description	Function	Did Item function properly (Yes/No)	Verified By	Date
Scale #1				
Power Button	With line power to the machine, does pushing the Power Button cause the Scale to start?			
Tare Button	With a weight on the scale, does pushing the Tare Button cause the scale reading to go to Zero?			
Scale #2				
Power Button	With line power to the machine, does pushing the Power Button cause the Scale to start?			
Tare Button	With a weight on the scale, does pushing the Tare Button cause the scale reading to go to Zero?			
Scale #3				
Power Button	With line power to the machine, does pushing the Power Button cause the Scale to start?			
Tare Button	With a weight on the scale, does pushing the Tare Button cause the scale reading to go to Zero?			
Scale #4				
Power Button	With line power to the machine, does pushing the Power Button cause the Scale to start?			
Tare Button	With a weight on the scale, does pushing the Tare Button cause the scale reading to go to Zero?			
Comments:				

Reviewed By: _____

Date: _____

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SCALES: CALIBRATION VERIFICATION



Equipment	Serial #	Calibration Date	Calibration Due Date	Verified By	Date
Multimeter					
Scale #1					
Scale #2					
Scale #3					
Scale #4					

Reviewed By: _____ Date: _____

ATTACHMENT - PROTOCOL DEVIATION REPORT LOG

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ATTACHMENT: PROTOCOL DEVIATION REPORT (PDR)

_____ General Information _____

System Name: _____ Protocol Number: _____

Deviation Report Number: _____ Protocol Step & Page No.: _____ -

_____ Instructions _____

1. The validation specialist assigns a sequential report number for each deviation with a specific protocol. For example, 001, 002, etc. can be easily referenced in a report.
2. Reference the relevant protocol number, step and page number of the noted deviation above.
3. Complete the below listed sections. If necessary, use additional pages and attach any supporting info.
4. Include the original PDR(s) with the protocol as an attachment. Summarize the impact of the deviation in the Validation Report.

Description of Deviation:

Investigation Evaluation and Results:

Corrective Action and Resolution:

Overall Investigation Review:

Prepared By: _____ Date: _____

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ATTACHMENT - SIGNATURE IDENTIFICATION LOG SHEET

Identify in the table below any personnel involved in the execution of this protocol.

Name	Affiliation	Signature	Initial	Date

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