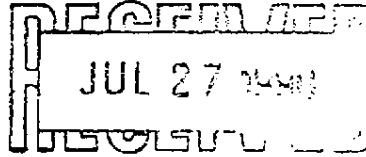


Vista Chemical Company

900 Threadneedle
Houston, Texas 77079
(713) 588-3000

P.O. Box 18028
Houston, Texas 77264
Fax (713) 588-3103

July 24, 1990



Mr. Joe Stratton
District Manager
VWR Scientific
P. O. Box 5025
Sugar Land, TX 77487

Dear Joe:

On behalf of Vista Chemical Company, I would like to express my sincere appreciation for providing our project team with the opportunity to tour your Westlake, LA facility and conduct an on-site quality evaluation survey.

Attached is a copy of the completed survey for your records. Your facility's overall rating of 82.4% ranks fairly high in our supplier evaluation program. Strengths were noted in most areas; however, I would like to mention the following items where we feel improvement is needed.

- Mission Statement - Your Mission Statement should be signed by senior management, CEO, etc.
- Certificates of Analysis - COA's should be readily available for chemicals stocked in your locations.
- Material Data Sheets - MDS's should be updated as necessary and sent to all Vista using locations, including Vista headquarters.
- Supplier Base - VWR's supplier base (approx. 1300) is extremely large. This could be reduced by eliminating marginal and less than marginal quality suppliers.

Joe, on the other side of the coin, if there is anything we (Vista) can do to assist your operations to run more smoothly, let's discuss at your earliest convenience.

I feel the communication link developed during our recent visit to your branch office will continue to grow and will assist the development of a long term relationship/partnership that will be mutually beneficial.

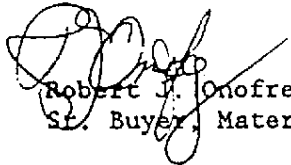
CWH 000012071

Mr. Joe Stratton
VWR Scientific
July 24, 1990

Page 2

If you have any questions or need additional information concerning this survey, please contact me at (713) 588-3307. Any comments you or your colleagues may have regarding this survey would certainly be appreciated. Thanks again for your cooperative and courteous treatment during our recent visit.

Sincerely,



Robert J. Onofrey
Sr. Buyer, Materials & MRO

RJO/ks
RJOWP51/RJ0020.90

CWH 000012072

bcc:

A. Bowman - LAB
B. Howell - VCM
B. McManemin - LCCP
M. Stillings - LCCP
W. H. Chamberlain
G. Draper
P. Markey
D. Hollis - LCCP
S. Vincent - LCCP
P. Fetzer - VCM
D. Whitehead - LCCP
A. Wallace - LAB
D. Phillips
F. Cooper - LCCP

DWH 000012073

VISTA SUPPLIER QUALITY EVALUATION

DETAILED SURVEY

SUPPLIER NAME VWR Scientific
ADDRESS 804 Columbia Southern Rd.
CITY, STATE, ZIP Westlake, LA 70669

SURVEY DATE 7-19-90 LAST SURVEY DATE _____

REASON FOR SURVEY: ☒ ROUTINE () FOLLOW-UP OR CORRECTIVE ACTION

IF FOLLOW-UP OR CORRECTIVE ACTION, PLEASE DESCRIBE: _____

PLEASE LIST SPECIFIC PRODUCTS SUPPLIED TO VISTA FROM THIS LOCATION:

<u>Lab Equipment and</u>	_____	_____
<u>Supplies</u>	_____	_____
_____	_____	_____
_____	_____	_____

SUPPLIER PERSONNEL CONTACTED DURING SURVEY

NAME	TITLE
1. <u>Joe Stratton</u>	<u>District Manager</u>
2. <u>Mike Arties</u>	<u>Distribution Manager</u>
3. <u>Zigmond Werlla</u>	<u>Lake Charles Office Manager</u>
4. <u>John Caillaver</u>	<u>Sales Representative</u>
5. _____	_____
6. _____	_____

WPI/DRP002

CWH 000012074

VISTA PERSONNEL PRESENT DURING SURVEY

NAME	TITLE
1. <u>Bob Onofrey</u>	<u>Sr. Buyer - Houston</u>
2. <u>Darwin Phillips</u>	<u>Buyer - Houston</u>
3. <u>Anne Bowman</u>	<u>Supervisor - Warehouse & Purchasing - LCLAS</u>
4. <u>Mike Stillings</u>	<u>Supervisor - Warehouse & Purchasing - LCLAS</u>
5. <u>Adrian Wallace</u>	<u>Chief Chemist - LCLAS</u>
6. <u>Paul Ferzer</u>	<u>Chief Chemist - LCVCM</u>
7. <u>Fred Cooke</u>	<u>Laboratory Supervisor - LCLAS</u>

IF VISTA HAS QUESTIONS CONCERNING IMPLEMENTATION OR INTERPRETATION OF THE STANDARDS AT THIS FACILITY, WHO SHOULD BE CONTACTED?

NAME Zigmond Werlla PHONE (318) 882-8000

SUPPLIER RATING RESULTS

	TOTAL POINTS	POINTS AWARDED	PERCENT OF POSSIBLE POINTS BY CATEGORY	CATEGORY WEIGHT FACTOR	POINTS AWARDED FOR CATEGORY
A. QUALITY MANAGEMENT PHILOSOPHY	<u>42</u>	<u>32</u>	<u>76.2%</u>	X 25%	<u>19.1</u>
B. MATERIAL CONTROL	<u>48</u>	<u>33</u>	<u>68.8%</u>	X 25%	<u>17.2</u>
C. INTERNAL QUALITY ASSURANCE	<u>48</u>	<u>36</u>	<u>75.0%</u>	X 25%	<u>18.8</u>
D. ORDER PROCESSING	<u>36</u>	<u>36</u>	<u>100%</u>	X 25%	<u>25.0</u>
TOTAL				100%	<u>82.4</u>

WP1/DRP002

CWH 000012075

VISTA SUPPLIER QUALITY EVALUATION

DETAILED SHEET

Review the following questions and evaluate the level of conformance based on the following rating:

Yes, full compliance - 2; partial compliance - 1; No - 0. ~~Where there are to be~~ considered: "Is there a system in effect?" (S), "Are there written procedures?" (P), "Is there documented compliance?" (C)

If there are any comments, indicate by checking the "See Comments" box (See Comm), and summarize any comments on an attachment sheet, referencing the question number.

If a question does not apply to the process, indicate by checking the N/A section.

A. QUALITY MANAGEMENT PHILOSOPHY

- | | 2 | 1 | 0 | See Comm | |
|--|-------|-------|-------|----------|---|
| 1. Is there a formal Company Quality Policy? | (2) | (1) | (0) | () | 0 |
| a) Does it have full support/involvement of top management? Mission Statement should have QA Management approval | (2) | (2) | (0) | () | 0 |
| b) Is there a documented Quality System (i.e., Quality Manual) and is it current? | (1) | (1) | (1) | () | 0 |
| 2. Is there evidence of commitment for continual quality improvement in all aspects of the business? | (2) | (2) | (2) | () | 0 |
| 3. Is there an ongoing program to assure that all employees are formally trained and involved in your quality procedures, policies and techniques? | (2) | (2) | (2) | () | 0 |
| 4. Is there a formal program to analyze and reduce customer complaints? | (2) | (2) | (2) | () | 0 |
| 5. Is there an internal audit system that evaluates the performance of the quality system? | (0) | (0) | (0) | () | 0 |

WP1/DRP002

CWH 000012076

3. MATERIAL CONTROL

- | | | | | | | | |
|-----|--|-------------------|-----|-----|-----|-----|---|
| 1. | Do you have specifications for all purchased material? | (2) | (2) | (2) | (2) | (2) | 6 |
| 2. | Do you maintain documentation and/or certification control on those materials you stock? | (1) | (1) | (1) | (1) | (1) | 3 |
| | a) Are they available to appropriate personnel (salesman, office, etc)? | Yes, your request | | | | | |
| 3. | Do you have a formal quality evaluation and rating system for your suppliers? | (2) | (1) | (1) | (1) | (1) | 5 |
| | a) Where is the evaluation conducted? | <u>On - Site</u> | | | | | |
| 4. | Do you verify incoming material quality against documented specifications? | (2) | (2) | (2) | (2) | (2) | 9 |
| 5. | Do you have a sampling and testing procedure for purchased material? | (0) | (0) | (1) | (1) | (1) | 3 |
| 6. | Is there a formal procedure for identification and segregation of non-conforming materials? | (2) | (2) | (2) | (2) | (2) | 6 |
| 7A. | Are your suppliers required to be involved in a QMP which utilizes statistical methods? | (1) | (2) | (2) | (1) | (1) | 4 |
| 7B. | If so, do you require your suppliers to submit SPC charts (data) to demonstrate process control? | () | () | () | (X) | () | 1 |
| 8. | How do you determine adequate inventory levels? | (2) | (2) | (2) | () | () | 6 |

WP1/DRP002

CWH 000012077

C. INTERNAL QUALITY ASSURANCE

- | | | | | | | | |
|----|---|---------------------|-----|-----|-----|-----|-----|
| 1. | Do you analyze non-conforming product returned by customers to: | | | | | | |
| | a) Determine failure mode? | (2) | (2) | (2) | (2) | (2) | (2) |
| | b) Develop action steps? | (2) | (2) | (2) | (2) | (2) | (2) |
| | c) Is a follow-up made with the customer? | (2) | (2) | (2) | (2) | (2) | (2) |
| 2. | Is material certification available to Vista, if required? | (2) | (2) | (2) | (2) | (2) | (2) |
| 3. | Do you have a system in place to monitor and use customer feedback? | (2) | (2) | (2) | (2) | (2) | (2) |
| 4. | Is the team concept utilized in problem solving? | (2) | (2) | (2) | (2) | (2) | (2) |
| 5. | Do you audit your quality program progress on a regular basis? | (0) | (0) | (0) | (0) | (0) | (0) |
| 6. | Is Quality Control a separate/distinct function in your organization? | (0) | (0) | (0) | (0) | (0) | (0) |
| | a) Who is responsible for Quality Control? | <u>Joe Stuckton</u> | | | | | |

D. ORDER PROCESSING

- | | | | | | | | |
|----|---|-------|-------|-------|-------|-------|---|
| 1. | What ensures orders are processed promptly and delivered on agreed date? | (2) | (2) | (2) | (2) | (2) | 6 |
| 2. | How do you trace and advise Vista on backorders? | (2) | (2) | (2) | (2) | (2) | 6 |
| 3. | Is there a system in effect to advise Vista of change in delivery schedules? | (2) | (2) | (2) | (2) | (2) | 6 |
| 4. | How do you ensure that Vista's terms, instructions (invoicing, destination, freight, etc) and pricing are reviewed before shipping and invoicing? | (2) | (2) | (2) | (2) | (2) | 6 |
| 5. | Is Vista advised of minimum order requirements? | () | () | () | (X) | () | |
| 6. | How do you verify <u>correct</u> material is being shipped? | (2) | (2) | (2) | () | () | 6 |
| 7. | How do you ensure adequate distribution of current product information? | (2) | (2) | (2) | () | () | 6 |

WP1/DRP002

CWH 000012079

E. VISUAL OBSERVATION

1. General Housekeeping Conditions:

- a) Clean Very
b) Needs Improvement

2. Condition of Equipment:

N/A

Excellent Good Fair Poor
1 2 3 4

3. Personnel

a) Helpfulness	<u>X</u>	<u></u>	<u></u>	<u></u>
b) Knowledgeable	<u>X</u>	<u></u>	<u></u>	<u></u>
c) Courteous	<u>X</u>	<u></u>	<u></u>	<u></u>
d) Availability	<u></u>	<u>X</u>	<u></u>	<u></u>
e) Reliability	<u>X</u>	<u></u>	<u></u>	<u></u>

WP1/DRP002

CWH 000012080

Vista Supplier Audit Form



ATTN:
Steve Smith
2

P.O. BOX 227
GEISMAR, LOUISIANA 70734
(504) 473-5000

A DIVISION OF
VULCAN MATERIALS COMPANY

CERTIFICATE OF ANALYSIS

For

Vista Chemical

CARBON TETRACHLORIDE

Customer Order No.:
Date Shipped: 12/20/90

Vulcan Order No.: 40316
Quantity: 77515
Lot No.: ST-40

COMPONENT

ANALYSIS

Color, APHA-----	3
Appearance-----	Clear
Specific Gravity @ 25/25°C-----	1.5890
Water, ppm-----	10
Acidity as HCL, ppm-----	0
Free Halogens-----	NONE

cc: Technical Service Department
VULCAN CHEMICALS
P.O. Box 7689
Birmingham, AL 35253-0689

Day Joyner
Supplier's Representative

Form No.: 540 1288

CWH 000012082

To _____

From _____

Calculation Sheet

VISTA

Audit Phases

Phase I

How do you ensure that we are receiving product that meets conforms to our specifications.

- Key Areas :- Sampling Frequency
- Items sampled
 - In-process indicators
 - Out of spec plans (corrective action)
 - Analytical methods - lab + On-line
- Accuracy
Precision
Consistency
Calibration

Phase II

How do you ensure that we are receiving consistent product?

- Key Areas - same as Phase I, Also
- SPC charts on Finished product
 - SPC on in process indicators
 - Frequency of off spec

Phase III

How do you ensure continual improvement in the product we are receiving?

- Key Areas - Phase I + II
- Focus on improvement areas.

CWH 000012083

Made By _____

Date _____

Page _____ of _____

Job No. _____

Title _____

VISTA SUPPLIER QUALITY EVALUATIONDETAILED SURVEY

Review the following questions and evaluate the level of conformance based on the following rating:

Yes, full compliance = 2; partial compliance = 1; No = 0. Three items are to be considered: "Is there a system in effect?" (S), "Are there written procedures?" (P), "Is there documented compliance?" (C)

If there are any comments, indicate by checking the "See Comments" box (See Comm), and summarize any comments on an attachment sheet, referencing the question number.

If a question does not apply to the process, indicate by checking the N/A section.

A. QUALITY MANAGEMENT PHILOSOPHY

		S	P	C	N/A	SEE COMM
		-----	-----	-----	-----	-----
<u>III</u>	1. Is there a formal Company Quality Policy?	()	()		()	()
<u>III</u>	2. Is there a documented Quality system (i.e., Quality manual) and is it current?	()	()	()	()	()
<u>III</u>	3. Is there evidence of commitment for continual quality improvement in all aspects of the business?	()	()	()	()	()
<u>III</u>	4. Is there an on-going program to assure that <u>all</u> employees are formally trained in your Quality procedures, policies and techniques?	()	()	()	()	()
<u>III</u>	5. Is there a formal program to analyze and reduce customer complaints?	()	()	()	()	()
<u>III</u>	6. Is there an internal audit system that evaluates the performance of the Quality System?	()	()	()	()	()

B. RAW MATERIAL CONTROL

S	P	C	N/A	SEE COMM
---	---	---	-----	-------------

- | | | | | | | | |
|-----|-----|---|-----|-----|-----|-----|-----|
| III | 1. | For purchased material, is there a system that assures the material meets your requirements? | () | () | () | () | () |
| III | 2. | Are there specifications for all purchased materials? | () | () | () | () | () |
| III | 3. | For purchased material, is there a system to routinely review/monitor specifications? | () | () | () | () | () |
| III | 4. | Are these specifications accessible to appropriate personnel? (i.e. QC Lab, Purchasing, Operations) | () | () | () | () | () |
| III | 5. | Is there a sampling and testing procedure for raw materials? (N/A if SPC or SQC charts are available for all raw materials) | () | () | () | () | () |
| III | 6. | Are suppliers required to furnish certificates of analysis and are they accessible to appropriate personnel i.e. QC Lab, Purchasing, Operations? (N/A if SQC charts are required for all raw materials) | () | () | () | () | () |
| III | 7. | Is there a formal procedure for the disposition of non-conforming raw materials? | () | () | () | () | () |
| III | 8. | Are rejected raw materials identified and physically segregated to prevent use? | () | () | | () | () |
| III | 9. | Is there a formal supplier evaluation and rating system? | () | () | () | () | () |
| III | 10. | Are suppliers required to be involved in a QMP which utilizes SPC techniques? | () | () | () | () | () |
| III | 11. | Are suppliers required to produce on-going evidence that their processes are in statistical control? | () | () | () | () | () |

C. IN-PROCESS CONTROL

		S	P	C	N/A	SEE COMM

II	1. Are all processes identified using flow diagrams?	()	()	()	()	()
I	2. Are there procedures to identify the following:					
	A. Operating procedures?	()	()	()	()	()
	B. Critical operating parameters?	()	()	()	()	()
	C. Potential process adjustments?	()	()	()	()	()
	D. Critical Process Measurements?	()	()	()	()	()
—	3. How are batches defined in this process?	_____				
I	4. Are in-process and finished batches traceable to raw materials?	()	()	()	()	()
I	5. Are unusual process operations reported to Quality Control?	()	()	()	()	()
I	6. Are calibration requirements of all in-process instrumentation prioritized and routinely followed? (i.e. preventative maintenance schedule)	()	()	()	()	()
II	7. Are control charts maintained on all critical process parameters?	()	()	()	()	()
II	8. Are charted processes monitored routinely to determine if they are in statistical control?	()	()	()	()	()
I	9. Are rework procedures properly authorized?	()	()	()	()	()
I	10. Is there a procedure to identify and dispose of all experimental batches?	()	()	()	()	()
I	11. Is there a first-piece verification or other suitable procedure to insure valid production setups upon product change?	()	()	()	()	()
I	12. During processing, are characteristics inspected or tested to assure the material consistently meets specification? (N/A if SPC charts are used in this process)	()	()	()	()	()
I	13. Are control charts used to monitor variations in test equipment or test procedures?	()	()	()	()	()

D. FINISHED PRODUCT CONTROL

		S	P	C	N/A	SEE COMM
I	1.	Are there current specifications for all products?				
		()	()	()	()	()
I	2.	Are there current test methods for all products?				
		()	()	()	()	()
II	3.	Are control charts used to monitor finished product specifications? (N/A if adequate control charts are maintained on process parameters.)				
		()	()	()	()	()
I	4.	Are Vista's specifications readily available to appropriate personnel?				
		()	()	()	()	()
I	5.	Is there a policy for the disposition of non-conforming finished product?				
		()	()	()	()	()
I	6.	Are Vista's packaging and coding requirements maintained and followed?				
		()	()	()	()	()
I	7.	Are shipping containers and product packages inspected for cleanliness and integrity?				
		()	()	()	()	()
I	8.	Is a zero discrepancy acceptance criterion used for product sampling programs (i.e. waivers are not allowed)?				
		()	()	()	()	()

E. QUALITY ASSURANCE

S	P	C	N/A	SEE COMM
---	---	---	-----	-------------

I 1. Is certification of product conformance to specification supplied to Vista? (N/A if not previously requested) () () () () ()

II 2. Is a system in place to monitor and use customer feedback? () () () () ()

III 3. Do you analyze non-conforming product returned by customers to:

A. Determine failure mode? () () () () ()

B. Develop action steps? () () () () ()

C. Follow-up with the customer? () () () () ()

I 4. Is Quality Control a separate and distinct part of the plant organization? () () () () ()

Who is in charge of QC? _____

~~III~~ II 5. Is SPC training provided to all personnel? () () () () ()

III 6. Is the team concept utilized in problem solving? () () () () ()

F. DELIVERY RATING

The following formula is used to determine the Delivery Rating summarized on Page 2:

DELIVERY RATING FORMULA

$$\frac{B}{Q} \times 100 = \text{Possible points for on-time deliveries}$$

Q - Total number of shipments received

B - Total number of shipments received on-time

Carry the above rating to Page 2 to calculate the overall survey rating.

G. INVOICE RATING

The following formula is used to determine the Invoice Rating Summarized on Page 2:

INVOICE RATING FORMULA

$$\frac{C}{T} \times 100 = \text{Possible points for correct invoicing}$$

C - Number of correct invoices received

T - Total number of invoices received.

Carry the above rating to Page 2 to calculate the overall survey rating.

VISTA SUPPLIER QUALITY EVALUATIONDETAILED SURVEY

SUPPLIER NAME _____

ADDRESS _____

CITY, STATE, ZIP _____

SURVEY DATE _____

LAST SURVEY DATE _____

REASON FOR SURVEY: () ROUTINE () FOLLOW-UP OR CORRECTIVE ACTION

IF FOLLOW-UP OR CORRECTIVE ACTION, PLEASE DESCRIBE: _____

PLEASE LIST SPECIFIC PRODUCTS PRODUCED FOR VISTA AT THIS LOCATION:

SUPPLIER PERSONNEL CONTACTED DURING SURVEY

NAMETITLE

1.	_____	_____
2.	_____	_____
3.	_____	_____
4.	_____	_____
5.	_____	_____
6.	_____	_____
7.	_____	_____
8.	_____	_____

WP7/SURVEY

1

CWH 000012091

VISTA PERSONNEL PRESENT DURING SURVEY

	<u>NAME</u>	<u>TITLE</u>
1.	_____	_____
2.	_____	_____
3.	_____	_____
4.	_____	_____
5.	_____	_____
6.	_____	_____

IF VISTA HAS QUESTIONS CONCERNING IMPLEMENTATION OR INTERPRETATION OF SPC PARAMETERS AT THIS FACILITY, WHO SHOULD BE CONTACTED?

NAME _____ PHONE _____

SUPPLIER RATING RESULTS

	POINTS AWARDED	POINTS POSSIBLE (1)	% POSSIBLE POINTS BY CATEGORY	CATEGORY IMPACT FACTOR	PERCENT OF POINTS AWARDED PER CATEGORY
A. QUALITY MANAGEMENT PHILOSOPHY	_____	_____	_____	X 15%	_____
B. RAW MATERIAL CONTROL	_____	_____	_____	X 20%	_____
C. IN-PROCESS CONTROL	_____	_____	_____	X 25%	_____
D. FINISH PRODUCT CONTROL	_____	_____	_____	X 10%	_____
E. QUALITY ASSURANCE	_____	_____	_____	X 10%	_____
F. DELIVERY RATING	_____	_____	_____	X 15%	_____
G. INVOICE RATING	_____	_____	_____	X 5%	_____
TOTAL	_____	_____	_____	_____	_____

(1) Excludes questions marked N/A

VISTA SUPPLIER QUALITY EVALUATIONDETAILED SURVEY

Review the following questions and evaluate the level of conformance based on the following rating:

Yes, full compliance = 2; partial compliance = 1; No = 0. Three items are to be considered: "Is there a system in effect?" (S), "Are there written procedures?" (P), "Is there documented compliance?" (C)

If there are any comments, indicate by checking the "See Comments" box (See Comm), and summarize any comments on an attachment sheet, referencing the question number.

If a question does not apply to the process, indicate by checking the N/A section.

A. QUALITY MANAGEMENT PHILOSOPHY

	S	P	C	N/A	SEE COMM
	=====				
1. Is there a formal Company Quality Policy?	()	()		()	()
2. Is there a documented Quality system (i.e., Quality manual) and is it current?	()	()	()	()	()
3. Is there evidence of commitment for continual quality improvement in all aspects of the business?	()	()	()	()	()
4. Is there an on-going program to assure that <u>all</u> employees are formally trained in your Quality procedures, policies and techniques?	()	()	()	()	()
5. Is there a formal program to analyze and reduce customer complaints?	()	()	()	()	()
6. Is there an internal audit system that evaluates the performance of the Quality System?	()	()	()	()	()

B. RAW MATERIAL CONTROL

	S	P	C	N/A	SEE COMM
1. For purchased material, is there a system that assures the material meets your requirements?	()	()	()	()	()
2. Are there specifications for all purchased materials?	()	()	()	()	()
3. For purchased material, is there a system to routinely review/monitor specifications?	()	()	()	()	()
4. Are these specifications accessible to appropriate personnel? (i.e. QC Lab, Purchasing, Operations)	()	()	()	()	()
5. Is there a sampling and testing procedure for raw materials? (N/A if SPC or SQC charts are available for all raw materials)	()	()	()	()	()
6. Are suppliers required to furnish certificates of analysis and are they accessible to appropriate personnel i.e. QC Lab, Purchasing, Operations? (N/A if SQC charts are required for all raw materials)	()	()	()	()	()
7. Is there a formal procedure for the disposition of non-conforming raw materials?	()	()	()	()	()
8. Are rejected raw materials identified and physically segregated to prevent use?	()	()		()	()
9. Is there a formal supplier evaluation and rating system?	()	()	()	()	()
10. Are suppliers required to be involved in a QMP which utilizes SPC techniques?	()	()	()	()	()
11. Are suppliers required to produce on-going evidence that their processes are in statistical control?	()	()	()	()	()

C. IN-PROCESS CONTROL

	S	P	C	N/A	SEE COMM
1. Are all processes identified using flow diagrams?	()	()	()	()	()
2. Are there procedures to identify the following:					
A. Operating procedures?	()	()	()	()	()
B. Critical operating parameters?	()	()	()	()	()
C. Potential process adjustments?	()	()	()	()	()
D. Critical Process Measurements?	()	()	()	()	()
3. How are batches defined in this process?					
4. Are in-process and finished batches traceable to raw materials?	()	()	()	()	()
5. Are unusual process operations reported to Quality Control?	()	()	()	()	()
6. Are calibration requirements of all in-process instrumentation prioritized and routinely followed? (i.e. preventative maintenance schedule)	()	()	()	()	()
7. Are control charts maintained on all critical process parameters?	()	()	()	()	()
8. Are charted processes monitored routinely to determine if they are in statistical control?	()	()	()	()	()
9. Are rework procedures properly authorized?	()	()	()	()	()
10. Is there a procedure to identify and dispose of all experimental batches?	()	()	()	()	()
11. Is there a first-piece verification or other suitable procedure to insure valid production setups upon product change?	()	()	()	()	()
12. During processing, are characteristics inspected or tested to assure the material consistently meets specification? (N/A if SPC charts are used in this process)	()	()	()	()	()
13. Are control charts used to monitor variations in test equipment or test procedures?	()	()	()	()	()

D. FINISHED PRODUCT CONTROL

	S	P	C	N/A	SEE COMM
1. Are there current specifications for all products?	()	()	()	()	()
2. Are there current test methods for all products?	()	()	()	()	()
3. Are control charts used to monitor finished product specifications? (N/A if adequate control charts are maintained on process parameters.)	()	()	()	()	()
4. Are Vista's specifications readily available to appropriate personnel?	()	()	()	()	()
5. Is there a policy for the disposition of non-conforming finished product?	()	()	()	()	()
6. Are Vista's packaging and coding requirements maintained and followed?	()	()	()	()	()
7. Are shipping containers and product packages inspected for cleanliness and integrity?	()	()	()	()	()
8. Is a zero discrepancy acceptance criterion used for product sampling programs (i.e. waivers are not allowed)?	()	()	()	()	()

E. QUALITY ASSURANCE

	S	P	C	N/A	SEE COMM
1. Is certification of product conformance to specification supplied to Vista? (N/A if not previously requested)	()	()	()	()	()
2. Is a system in place to monitor and use customer feedback?	()	()	()	()	()
3. Do you analyze non-conforming product returned by customers to:					
A. Determine failure mode?	()	()	()	()	()
B. Develop action steps?	()	()	()	()	()
C. Follow-up with the customer?	()	()	()	()	()
4. Is Quality Control a separate and distinct part of the plant organization?	()	()	()	()	()
Who is in charge of QC?					
5. Is SPC training provided to all personnel?	()	()	()	()	()
6. Is the team concept utilized in problem solving?	()	()	()	()	()

F. DELIVERY RATING

The following formula is used to determine the Delivery Rating summarized on Page 2:

DELIVERY RATING FORMULA

$$\frac{B}{Q} \times 100 = \text{Possible points for on-time deliveries}$$

Q - Total number of shipments received

B - Total number of shipments received on-time

Carry the above rating to Page 2 to calculate the overall survey rating.

G. INVOICE RATING

The following formula is used to determine the Invoice Rating Summarized on Page 2:

INVOICE RATING FORMULA

$$\frac{C}{T} \times 100 = \text{Possible points for correct invoicing}$$

C = Number of correct invoices received

T = Total number of invoices received.

Carry the above rating to Page 2 to calculate the overall survey rating.

M. Kane
JCM

VISTA

Interoffice
Communication

TO: Distribution
FROM: Anne Bowman
DATE: April 22, 1988
SUBJECT: Minutes from QMP Vendor Evaluation
Meeting of April 19, 1988

Attendees: Bob Onofrey, Darwin Phillips, Mike Stillings,
Bill Howell, Anne Bowman

Anne Bowman was appointed secretary.

Bob Onofrey opened the meeting by stating the purpose of
this meeting was to finalize our Vendor Evaluation Form.

The second category, Material Control, was reviewed and
the following questions were agreed to be included.

B.) Material Control	S	P	C	N/A	See Comm.
1.) Do you have specifications for all purchased material?	()	()	()	()	()
2.) Do you maintain documentation and/or certification control on those materials you stock?	()	()	()	()	()
A.) Are they available to appropriate personnel (salesman, office, etc.)	()	()	()	()	()
3.) Do you have a formal quality evaluation and rating system for your suppliers?	()	()	()	()	()
A.) Where is the evaluation conducted? Mfg. Plant___ Other___	()	()	()	()	()
4.) Do you verify incoming materials quality against documented specifications?	()	()	()	()	()

	S	P	C	N/A	See Comm.
5.) Do you have a sampling and testing procedure for purchased material?	()	()	()	()	()
6.) Is there a formal procedure for identification and segregating of non-conforming materials?	()	()	()	()	()
7.) Are your suppliers required to be involved in a QMP which utilizes statistical methods?	()	()	()	()	()
8.) Is there a system to maintain adequate inventory levels?	()	()	()	()	()

Category #3, was originally to be "Order Processing/Invoicing" but it was agreed to be deleted because some of the same information could be obtained and included in the category of "Daily Performance".

The third category, Internal Quality Assurance, was reviewed and the following questions were agreed to be included:

C.) Internal Quality Assurance	S	P	C	N/A	See Comm.
1.) Do you analyze non-conforming product returned by customers to:	()	()	()	()	()
A. Determine failure mode?	()	()	()	()	()
B. Develop action steps?	()	()	()	()	()
C. Is a follow-up made with the customer?	()	()	()	()	()
2.) Is material certification available to Vista, if required?	()	()	()	()	()
3.) Do you have a system in place to monitor and use customer feedback?	()	()	()	()	()
4.) Is the team concept utilized in problem solving?	()	()	()	()	()

CWH 000012101

	S	P	C	N/A	See Comm.
6.) Is Quality Control a separate/distinct function in your organization?	()	()	()	()	()
A. Who is responsible for Quality Control?	()	()	()	()	()

The fourth category, Daily Performance, was reviewed and the following questions were agreed to be included:

D.) Daily Performance	S	P	C	N/A	See Comm.
1.) How do you ensure orders are processed promptly and delivered on agreed date?	()	()	()	()	()
2.) How do you trace and advise Vista on backorders?	()	()	()	()	()
3.) Is there a system in effect to advise Vista of change in delivery schedules?	()	()	()	()	()
4.) How do you ensure that Vista's terms, instructions (invoicing, destination, freight, etc.) and pricing are reviewed before shipping and invoicing?	()	()	()	()	()
5.) Is Vista advised of minimum order requirements?	()	()	()	()	()
6.) How do you verify <u>correct</u> material is being shipped?	()	()	()	()	()
7.) How do you insure adequate distribution of current product information?	()	()	()	()	()

The fifth category for general overall review, Visual Observation, was reviewed and the following was agreed to be included:

E.) Visual Observation

1.) General Housekeeping Conditions

- A. Clean _____
B. Needs Improvement _____

2.) Condition of equipment _____

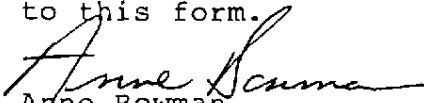
	<u>Excellent</u> 1	<u>Good</u> 2	<u>Poor</u> 3
3.) Personnel			
A. Helpfulness	_____	_____	_____
B. Knowledgeable	_____	_____	_____
C. Courteous	_____	_____	_____
D. Availability	_____	_____	_____
E. Reliability	_____	_____	_____

The next topic of discussion was the Vendors Rating Results. The following was decided.

	<u>Total</u> <u>Points(1)</u>	<u>Points</u> <u>Awarded</u>	<u>Percent of</u> <u>Possible Category</u> <u>Points By Impact</u> <u>Category Factor</u>	<u>Percentage</u> <u>Points</u> <u>Awarded Per</u> <u>Category</u>
A. Quality Management Philosophy	(42)	_____	% X 15%	_____
B. Material Contract	(48)	_____	% X 25%	_____
C. Internal Quality Assurance	(48)	_____	% X 25%	_____
D. Daily Performance	(42)	_____	% X 35%	_____
TOTAL	(180)	_____	100%	_____

(1) Total points possible less 6 points for each N/A.

Attached is a proof copy of the final form. The next meeting is scheduled for Thursday, May 19, 1988 at 8:00 a.m. at the LCLAB Plant to make any changes or corrections to this form.


Anne Bowman

Distribution: RJO, DRP - Houston Purchasing
MGS - LCCP
WRH - VCM

cc: W. H. McNeese - Houston
W. H. Chamberlain - Houston
R. D. Gamblin - Houston
T. F. Wall - LCLAB
P. R. Ardoin - LCCP
A. G. Loudon - LCLAB
M. R. Kane - LCVCM
P. E. Markey - LCVCM
G. G. Draper - Houston
B. H. Swan - LCCP

CWH 000012103

VISTA QMP MRO/DISTRIBUTOR SUPPLIER
QUALITY EVALUATION

DETAILED SURVEY

SUPPLIER NAME _____

ADDRESS _____

CITY, STATE, ZIP _____

SURVEY DATE _____ LAST SURVEY DATE _____

REASON FOR SURVEY: () ROUTINE () FOLLOW-UP OR CORRECTIVE
ACTION

IF FOLLOW-UP OR CORRECTIVE ACTION, PLEASE DESCRIBE: _____

PLEASE LIST SPECIFIC PRODUCTS PROVIDED TO VISTA AT THIS
LOCATION:

_____	_____	_____
_____	_____	_____
_____	_____	_____
_____	_____	_____

SUPPLIER PERSONNEL CONTACTED DURING SURVEY

	NAME	TITLE
1.	_____	_____
2.	_____	_____
3.	_____	_____
4.	_____	_____
5.	_____	_____
6.	_____	_____

WP1/DRP002

CWH 000012104

VISTA PERSONNEL PRESENT DURING SURVEY

	NAME	TITLE
1.	_____	_____
2.	_____	_____
3.	_____	_____
4.	_____	_____
5.	_____	_____
6.	_____	_____

IF VISTA HAS QUESTIONS CONCERNING IMPLEMENTATION OR INTERPRETATION OF SPC PARAMETERS AT THIS FACILITY, WHO SHOULD BE CONTACTED?

NAME _____ PHONE _____

	Total Points(1)	Points Awarded	Percent of Possible Category Points By Category	Impact Factor	Percentage Points Awarded Per Category
A. Quality Management Philosophy	(42)	_____	_____ %	X 15%	_____
B. Material Contract	(48)	_____	_____ %	X 25%	_____
C. Internal Quality Assurance	(48)	_____	_____ %	X 25%	_____
D. Daily Performance	(42)	_____	_____ %	X 35%	_____
 TOTAL	 (180)	 _____		 100%	 _____

(1) Total points possible less 6 points for each N/A.

VISTA QMP MRO/DISTRIBUTION SUPPLIER
QUALITY EVALUATION

DETAILED SURVEY

Review the following questions and evaluate the level of conformance based on the following rate:

Yes, full compliance = 2; partial compliance = 1; No = 0.
Three items are to be considered: "Is there a system in effect?" (S), "Are there written procedures?" (P), "Is there documented compliance?" (C)

If there are any comments, indicate by checking the "See Comments" box (See Comm.), and summarize any comments on an attachment sheet, referencing the question number.

	S	P	C	N/A	See Comm.
A.) Quality Management Philosophy					
1.) Is there a formal Company Quality Policy?	()	()	()	()	()
a.) Does it have full support/involvement of Top Management?	()	()	()	()	()
b.) Is there a documented Quality system (i.e. Quality Manual) and is it current?	()	()	()	()	()
2.) Is there evidence of commitment for continual quality improvement in all aspects of the business?	()	()	()	()	()
3.) Is there an ongoing program to assure that all employees are formally trained and involved in your quality procedures, policies, and techniques?	()	()	()	()	()
4.) Is there a formal program to analyze and reduce customer complaints?	()	()	()	()	()
5.) Is there an internal audit system that evaluates the performance of the quality system?	()	()	()	()	()

	S	P	C	N/A	See Comm.
B. Material Control					
1.) Do you have specifications for all purchased material?	()	()	()	()	()
2.) Do you maintain documentation and/or certification control on those materials you stock?	()	()	()	()	()
A. Are they available to appropriate personnel (salesman, office, etc.)?	()	()	()	()	()
3.) Do you have a formal quality evaluation and rating system for your suppliers?	()	()	()	()	()
4.) Do you verify incoming materials quality against documented specifications?	()	()	()	()	()
5.) Do you have a sampling and testing procedure for purchased material?	()	()	()	()	()
6.) Is there a formal procedure for identification and segregating of non-conforming materials?	()	()	()	()	()
7.) Are your suppliers required to be involved in a QMP which utilizes statistical methods?	()	()	()	()	()
8.) Is there a system to maintain adequate inventory levels?	()	()	()	()	()
C.) Internal Quality Assurance					
1.) Do you analyze non-conforming product returned by customers to:	()	()	()	()	()
A. Determine failure mode?	()	()	()	()	()
B. Develop action steps?	()	()	()	()	()
C. Is a follow-up made with the customer?	()	()	()	()	()
2.) Is material certification available to Vista, if required?	()	()	()	()	()

	S	P	C	N/A	See Comm.
3.) Do you have a system in place to monitor and use customer feedback?	()	()	()	()	()
4.) Is the team concept utilized in problem solving?	()	()	()	()	()
5.) Do you audit your quality program progress on a regular basis?	()	()	()	()	()
6.) Is Quality Control a separate/distinct function in your organization?	()	()	()	()	()

D.) Daily Performance

1.) How do you ensure orders are processed promptly and delivered on agreed date?	()	()	()	()	()
2.) How do you trace and advise Vista on backorders?	()	()	()	()	()
3.) Is there a system in effect to advise Vista of change in delivery schedules?	()	()	()	()	()
4.) How do you ensure that Vista's terms, instructions (invoicing, destination, freight, etc.) and pricing are reviewed before shipping and invoicing?	()	()	()	()	()
5.) Is Vista advised of minimum order requirements?	()	()	()	()	()
6.) How do you verify <u>correct</u> material is being shipped?	()	()	()	()	()
7.) How do you insure adequate distribution of current product information?	()	()	()	()	()

E.) Visual Observation

1.) General Housekeeping Conditions

- A. Clean _____
 B. Needs Improvement _____

CWH 000012108

2.) Condition of Equipment _____

<u>Excellent</u>	<u>Good</u>	<u>Poor</u>
<u>1</u>	<u>2</u>	<u>3</u>

3.) Personnel

A. Helpfulness	_____	_____	_____
B. Knowledgeable	_____	_____	_____
C. Courteous	_____	_____	_____
D. Availability	_____	_____	_____
E. Reliability	_____	_____	_____

VISTA SUPPLIER QUALITY EVALUATION

DETAILED SURVEY

SUPPLIER NAME _____

ADDRESS _____

CITY, STATE, ZIP _____

SURVEY DATE _____ LAST SURVEY DATE _____

REASON FOR SURVEY: () ROUTINE () FOLLOW-UP OR CORRECTIVE ACTION

IF FOLLOW-UP OR CORRECTIVE ACTION, PLEASE DESCRIBE: _____

PLEASE LIST SPECIFIC PRODUCTS PRODUCED FOR VISTA AT THIS LOCATION:

_____	_____	_____
_____	_____	_____
_____	_____	_____
_____	_____	_____

SUPPLIER PERSONNEL CONTACTED DURING SURVEY

	<u>NAME</u>	<u>TITLE</u>
1.	_____	_____
2.	_____	_____
3.	_____	_____
4.	_____	_____
5.	_____	_____
6.	_____	_____

WP1/DRP002

CWH 000012110

VISTA PERSONNEL PRESENT DURING SURVEY

	<u>NAME</u>	<u>TITLE</u>
1.	_____	_____
2.	_____	_____
3.	_____	_____
4.	_____	_____
5.	_____	_____
6.	_____	_____

IF VISTA HAS QUESTIONS CONCERNING IMPLEMENTATION OR INTERPRETATION OF SPC PARAMETERS AT THIS FACILITY, WHO SHOULD BE CONTACTED?

NAME _____ PHONE _____

PLEASE INDICATE THE GENERAL CONDITION OF THE FACILITIES, HOUSEKEEPING, ETC.

SUPPLIER RATING RESULTS

	POINTS AWARDED	POINTS POSSIBLE (1)	PERCENT OF POSSIBLE POINTS BY CATEGORY	CATEGORY IMPACT FACTOR	PERCENTAGE POINTS AWARDED PER CATEGORY
A. QUALITY MANAGEMENT PHILOSOPHY	_____	_____	_____ %	X 15%	_____
B. RAW MATERIAL CONTROL	_____	_____	_____ %	X 20%	_____
C. IN-PROCESS CONTROL	_____	_____	_____ %	X 30%	_____
D. FINISH PRODUCT CONTROL	_____	_____	_____ %	X 20%	_____
E. QUALITY ASSURANCE	_____	_____	_____ %	X 15%	_____
TOTALS	_____	_____			_____

(1) TOTAL POINTS POSSIBLE LESS 6 POINTS FOR EACH N/A
WP1/DRP002

CWH 000012111

VISTA SUPPLIER QUALITY EVALUATION

DETAILED SURVEY

Review the following questions and evaluate the level of conformance based on the following rating:

Yes, full compliance = 2; partial compliance = 1; No = 0. Three items are to be considered: "Is there a system in effect?" (S), "Are there written procedures?" (P), "Is there documented compliance?" (C)

If there are any comments, indicate by checking the "See Comments" box (See Comm), and summarize any comments on an attachment sheet, referencing the question number.

If a question does not apply to the process, indicate by checking the N/A section.

A. QUALITY MANAGEMENT PHILOSOPHY

- | | S | P | C | N/A | SEE
COMM |
|---|-----|-----|-----|-----|-------------|
| 1. Is there a formal Company Quality Policy? | () | () | () | () | () |
| Is there a documented Quality system (i.e., Quality manual) and is it current? | () | () | () | () | () |
| 2. Is there evidence of commitment for continual quality improvement in all aspects of the business? | () | () | () | () | () |
| 3. Is there an on-going program to assure that <u>all</u> employees are formally trained in your Quality procedures, policies and techniques? | () | () | () | () | () |
| 4. Is there a formal program to analyze and reduce customer complaints? | () | () | () | () | () |
| 5. Is there an internal audit system that evaluates the performance of the Quality System? | () | () | () | () | () |

B. RAW MATERIAL CONTROL

- | | S | P | C | N/A | SEE
COMM |
|--|-----|-----|-----|-----|-------------|
| 1. For purchased material is there a system that assures the material meets your requirements? | () | () | () | () | () |
| 2. Are there specifications for all purchased materials? | () | () | () | () | () |

WP1/DRP002

CWH 000012112

	S	P	C	N/A	SEE COMM
3. For purchased material, is there a system to review/monitor specifications?	()	()	()	()	()
4. Are these specifications accessible to appropriate personnel? (i.e. QC Lab, Purchasing, Operations)	()	()	()	()	()
5. Is there a sampling and testing procedure for raw materials?	()	()	()	()	()
6. Are suppliers required to furnish certificates of analysis and are they accessible to appropriate personnel? (i.e. QC Lab, Purchasing, Operations)	()	()	()	()	()
7. Is there a formal procedure for the disposition of non-conforming raw materials?	()	()	()	()	()
8. Are rejected raw materials identified and physically segregated to prevent use?	()	()	()	()	()
9. Is there a formal vendor evaluation and rating system?	()	()	()	()	()
10. Are suppliers required to be involved in a QMP which utilizes SPC techniques?	()	()	()	()	()
11. Are suppliers required to produce on-going evidence that their processes are in statistical control?	()	()	()	()	()

C. IN-PROCESS CONTROL

	S	P	C	N/A	SEE COMM
1. Are all processes identified using flow diagrams? Attach all pertinent PFD's.	()	()	()	()	()
2. Are there procedures to identify the following:					
A. Operating procedures?	()	()	()	()	()
B. Critical parameters?	()	()	()	()	()
C. Potential Process adjustments?	()	()	()	()	()
D. Process Measurements?	()	()	()	()	()

	S	P	C	N/A	SEE COMM
3. Are in-process and finished batches traceable to raw materials? How are batches defined in this process?	()	()	()	()	()
4. Are unusual process operations reported to Quality Control?	()	()	()	()	()
5. Are calibration requirements of all instrumentation prioritized and routinely followed?	()	()	()	()	()
6. Is there a formal preventive maintenance schedule?	()	()	()	()	()
7. Are control charts maintained on all critical process parameters?	()	()	()	()	()
8. Are charted processes monitored routinely to determine if they are in statistical control?	()	()	()	()	()
9. Are rework procedures properly authorized?	()	()	()	()	()
10. Is there a procedure to identify and dispose of all experimental batches?	()	()	()	()	()
11. Is there a first-piece verification or other suitable procedure to insure valid production setups?	()	()	()	()	()
12. During processing, are characteristics inspected or tested to assure the material consistently meets specification?	()	()	()	()	()

D. FINISHED PRODUCT CONTROL

	S	P	C	N/A	SEE COMM
1. Are there specifications:					
a. For all products?	()	()	()	()	()
b. Are they current?	()	()	()	()	()
2. Are there test methods:					
a. For all products?	()	()	()	()	()
b. Are they current?	()	()	()	()	()

	S	P	C	N/A	SEE COMM
3. Are control charts used to monitor finished product specifications?	()	()	()	()	()
4. Are Vista's specifications readily available to appropriate personnel?	()	()	()	()	()
5. Is there a policy for the disposition of non-conforming finished product?	()	()	()	()	()
6. Are Vista's packaging and coding requirements maintained and followed?	()	()	()	()	()
7. Are shipping containers and product packages inspected for cleanliness and integrity?	()	()	()	()	()
8. Is a zero discrepancy acceptance criterion used for product sampling programs?	()	()	()	()	()

E. QUALITY ASSURANCE

	S	P	C	N/A	SEE COMM
1. Is there a Quality assurance manual defining the Quality control operations and functions for specific processes?	()	()	()	()	()
2. Is there traceability of the delivered product back to raw materials?	()	()	()	()	()
3. Is certification of product conformance to specification supplied to Vista?	()	()	()	()	()
4. Do you analyze non-conforming product returned by customers to:					
A. determine failure mode?	()	()	()	()	()
B. Develop action steps?	()	()	()	()	()
C. Is a follow-up made with the customer?	()	()	()	()	()
5. Is Quality Control a separate and distinct part of the plant organization? Who is in charge of QC?	()	()	()	()	()
6. Has statistical process control been implemented in this facility?	()	()	()	()	()

- | | S | P | C | N/A | SEE
COMM |
|--|-----|-----|-----|-----|-------------|
| 7. Is SPC training provided to all personnel? | () | () | () | () | () |
| 8. Is the team concept utilized in problem solving? | () | () | () | () | () |
| 9. Are control charts used to monitor variations in test equipment? | () | () | () | () | () |
| 10. Is a system in place to monitor and use customer feedback? | () | () | () | () | () |
| 11. Attach a general block flow diagram of your order entry process through invoice mailing. | | | | | |

WP1/DRP002

CWH 000012116

Interoffice
Communication

To: Distribution

From: D. R. Phillips
Date: May 31, 1988

Subject: SUPPLIER QUALITY EVALUATION FORM

VISTA

The attached Supplier Quality Evaluation Form for MRO Suppliers and Distributors was developed as a tool to compliment our existing detailed survey for manufacturers.

The team would appreciate your comments on the form by Wednesday, June 15.

Thanks.



D. R. Phillips

DRP/rr

Distribution:	P. R. Ardoin, LCCP	E. McNinch, Baltimore
	D. A. Barclay, Blane	J. J. McLaughlin, OKC
	L. R. Bauer, Baltimore	W. H. McNeese
	W. H. Chamberlain	C. R. Miller, Blane
	R. A. Conrad, VCM	A. E. Russell, Hammond
	J. A. DeBernardi	S. K. Saborsky, Premiere
	R. G. DeYoung, Premiere	R. J. Schettek
	J. Friend, LCCP	R. W. Seymour, Aberdeen
	R. D. Gamblin	A. W. Sirmons, OKC
	H. D. Garrison, OKC	B. H. Swan, LCCP
	D. W. Hollis, Aberdeen	M. A. Swierc, Aberdeen
	T. H. Huffman	M. F. Ticar, Baltimore
	J. E. Little, Aberdeen	A. T. Trent
	M. R. Kane, VCM	T. F. Wall, LAB
	A. G. Loudon, LAB	J. W. Ware, LAB
	J. B. Maher, Hammond	C. C. Wyche, Baltimore
	P. E. Markey, VCM	

WP7/DRP025

CWH 000012117

VISTA SUPPLIER QUALITY EVALUATION

DETAILED SURVEY

SUPPLIER NAME _____

ADDRESS _____

CITY, STATE, ZIP _____

SURVEY DATE _____

LAST SURVEY DATE _____

REASON FOR SURVEY: () ROUTINE () FOLLOW-UP OR CORRECTIVE ACTION

IF FOLLOW-UP OR CORRECTIVE ACTION, PLEASE DESCRIBE: _____

PLEASE LIST SPECIFIC PRODUCTS PRODUCED FOR VISTA AT THIS LOCATION:

SUPPLIER PERSONNEL CONTACTED DURING SURVEY

NAME

TITLE

1. _____

2. _____

3. _____

4. _____

5. _____

6. _____

CWH 000012118

VISTA PERSONNEL PRESENT DURING SURVEY

	<u>NAME</u>	<u>TITLE</u>
1.	_____	_____
2.	_____	_____
3.	_____	_____
4.	_____	_____
5.	_____	_____
6.	_____	_____

IF VISTA HAS QUESTIONS CONCERNING IMPLEMENTATION OR INTERPRETATION OF SPC PARAMETERS AT THIS FACILITY, WHO SHOULD BE CONTACTED?

NAME _____ PHONE _____

SUPPLIER RATING RESULTS

	TOTAL POINTS	POINTS AWARDED	PERCENT OF POSSIBLE POINTS BY CATEGORY	CATEGORY IMPACT FACTOR	PERCENTAGE POINTS AWARDED PER CATEGORY
A. QUALITY MANAGEMENT PHILOSOPHY	_____	_____	_____ %	X 15%	_____
B. MATERIAL CONTROL	_____	_____	_____ %	X 25%	_____
C. INTERNAL QUALITY ASSURANCE	_____	_____	_____ %	X 25%	_____
D. ORDER PROCESSING	_____	_____	_____ %	X 35%	_____
TOTAL				100%	

VISTA SUPPLIER QUALITY EVALUATION

DETAILED SURVEY

Review the following questions and evaluate the level of conformance based on the following rating:

Yes, full compliance = 2; partial compliance = 1; No = 0. Three items are to be considered: "Is there a system in effect?" (S), "Are there written procedures?" (P), "Is there documented compliance?" (C)

If there are any comments, indicate by checking the "See Comments" box (See Comm), and summarize any comments on an attachment sheet, referencing the question number.

If a question does not apply to the process, indicate by checking the N/A section.

A. QUALITY MANAGEMENT PHILOSOPHY

	S	P	C	N/A	SEE COMM
1. Is there a formal Company Quality Policy?	()	()	()	()	()
a) Does it have full support/involvement of top management?	()	()	()	()	()
b) Is there a documented Quality System (i.e., Quality Manual) and is it current?	()	()	()	()	()
2. Is there evidence of commitment for continual quality improvement in all aspects of the business?	()	()	()	()	()
3. Is there an ongoing program to assure that all employees are formally trained and involved in your quality procedures, policies and techniques?	()	()	()	()	()
4. Is there a formal program to analyze and reduce customer complaints?	()	()	()	()	()
5. Is there an internal audit system that evaluates the performance of the quality system?	()	()	()	()	()

WP1/DRP002

CWH 000012120

B. MATERIAL CONTROL

	S	P	C	N/A	SEE COMM
1. Do you have specifications for all purchased material?	()	()	()	()	()
2. Do you maintain documentation and/or certification control on those materials you stock?	()	()	()	()	()
a) Are they available to appropriate personnel (salesman, office, etc)?					
3. Do you have a formal quality evaluation and rating system for your suppliers?	()	()	()	()	()
a) Where is the evaluation conducted?					
4. Do you verify incoming material quality against documented specifications?	()	()	()	()	()
5. Do you have a sampling and testing procedure for purchased material?	()	()	()	()	()
6. Is there a formal procedure for identification and segregation of non-conforming materials?	()	()	()	()	()
7. Are your suppliers required to be involved in a QMP which utilizes statistical methods?	()	()	()	()	()
8. How do you determine adequate inventory levels?	()	()	()	()	()

C. INTERNAL QUALITY ASSURANCE

S P C N/A SEE
COMM

1. Do you analyze non-conforming product returned by customers to:
 - a) Determine failure mode? () () () () ()
 - b) Develop action steps? () () () () ()
 - c) Is a follow-up made with the customer? () () () () ()
2. Is material certification available to Vista, if required? () () () () ()
3. Do you have a system in place to monitor and use customer feedback? () () () () ()
4. Is the team concept utilized in problem solving? () () () () ()
5. Do you audit your quality program progress on a regular basis? () () () () ()
6. Is Quality Control a separate/distinct function in your organization? () () () () ()
 - a) Who is responsible for Quality Control? _____

D. ORDER PROCESSING

	S	P	C	N/A	SEE COMM
1. What ensures orders are processed promptly and delivered on agreed date?	()	()	()	()	()
2. How do you test and advise Vista on backorders?	()	()	()	()	()
3. Is there a system in effect to advise Vista of change in delivery schedules?	()	()	()	()	()
4. How do you ensure that Vista's terms, instructions (invoicing, destination, freight, etc) and pricing are reviewed before shipping and invoicing?	()	()	()	()	()
5. Is Vista advised of minimum order requirements?	()	()	()	()	()
6. How do you verify <u>correct</u> material is being shipped?	()	()	()	()	()
7. How do you ensure adequate distribution of current product information?	()	()	()	()	()

E. VISUAL OBSERVATION

1. General Housekeeping Conditions:

- a) Clean _____
b) Needs Improvement _____

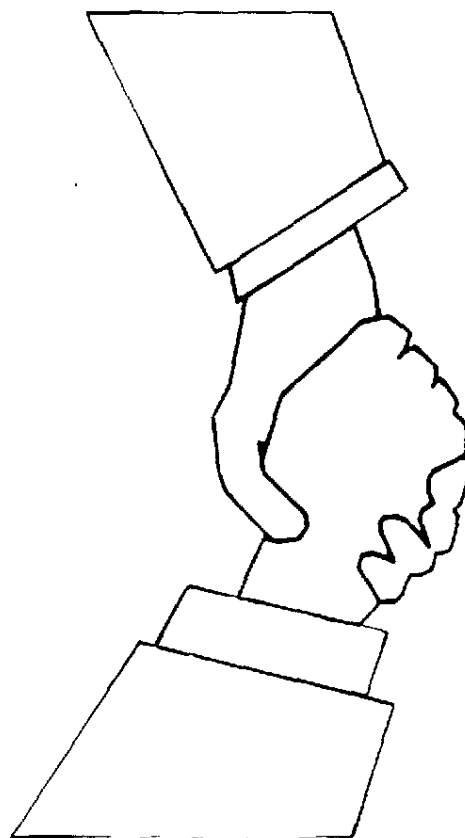
2. Condition of Equipment:

<u>Excellent</u>	<u>Good</u>	<u>Poor</u>	<u>See Comm</u>
1	2	3	

3. Personnel

a) Helpfulness	_____	_____	_____	_____
b) Knowledgeable	_____	_____	_____	_____
c) Courteous	_____	_____	_____	_____
d) Availability	_____	_____	_____	_____
e) Reliability	_____	_____	_____	_____

SUPPLIER PARTNERSHIP DEVELOPMENT



CWH 000012125

NEW MATERIALS MANAGEMENT PHILOSOPHY

PARTNERSHIPS



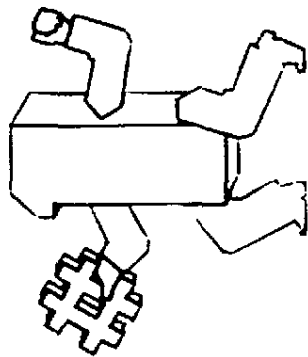
Trust
Single-sourcing
Early supplier involvement
Design of Experiments
Commodity Teams
Just-In-Time deliveries

Table 4. Why a supplier-customer partnership is a win-win relationship.

<i>Buyer Responsibilities- Supplier Benefits</i>	<i>Mutual Responsibilities</i>	<i>Supplier Responsibilities- Buyer Benefits</i>
• Longer-term contracts • Larger volume • Fewer suppliers • Consultation in design • Technical help • Quality assistance • Higher yields • More stable forecasts • Fair dealings, fair profit, fair ROI	• Trust, loyalty • Long-term marriage • Win-win relationship • Mutually acceptable specifications • Financial incentives/penalties	• Excellent quality • Lower prices • Early delivery • Reduced cycle time • Ideas for improvement • Design assistance • Standardization assistance

TABLE A: THE FOUR STAGES OF SUPPLY MANAGEMENT

AREA	STAGE 1 CONFRONTATION	STAGE 2 ARMS LENGTH	STAGE 3 GOAL CONGRUENCE	STAGE 4 FULL PARTNERSHIP
Supplier Relationships	• Distrust	• Suspicion	• Limited Trust	• Full Trust
	• Supplier Proliferation	• Multiple Suppliers	• Reduced Supplier Base	• Single Sourcing
	• Economic Domination	• Laissez Faire	• Preferred Suppliers	• Partnership Suppliers
	• Supplier Kept in Dark	• Limited Information	• Mutual Consultations	• Frequent Visits, Training, and Technical Help
Management	• DL Focus	• Automation Focus	• FMS Focus	• Supply Mgmt. Focus
	• Purchasing - An Appendage	• Cost Pre-occupation	• Quality Pre-occupation	• Emphasis on Quality, Cost and Cycle Time
	• Churn Mgmt.	• Nomadic Mgmt.	• Settled Mgmt.	• Team Management
	• No Commitment to Partnership	• Cautious Overtures	• "Open Kimono" Policy	• Early Supplier Involvement
	• No Tapping or Supplier Creativity	• Creativity Neutrality	• Supplier Cost Reductions with No Sharing	• Savings Sharing
Organization	• Functional Barriers	• Matrix	• Centralized Supply Mgmt.	• Commodity Teams
	• Purchasing De-centralized	• Sourcing - Purchasing Separation	• SOA - Part of Supply Mgmt.	• ENG, SOA, Purchasing and Supplier One Team
Measurement	• Price, Price, Price	• Stock-outs	• Inventory Turns	• Total Cost (Price and Cost of Poor Quality, Delivery)
	• Variance from Budget	• Line Shut-downs	• Lead Time Reduction	• Total Cycle Time
Quality	• Specs: Vague, Arbitrary	• Specs: Formula Computer Approximate	• Classification of Characteristics	• Specs Thru Q.F.D.
	• Boiler Plate A.Q.L.s	• Reduced A.Q.L.s	• Zero Defects	• Target Values
	• Heavy Incoming Insp.	• Reduced Incoming Insp.	• Skip Lot	• Certification
	• C_{pk} Unknown	• $C_{pk} > 1.33$	• $C_{pk} > 2.0$	• $C_{pk} > 5.0$
	• S.P.C. Unknown	• S.P.C. = Control Charts	• D.O.E. to Solve Problems & Pre-control for S.P.C.	• D.O.E. at Design Stage of Product/Process
Cost	• 3 Quote Syndrome	• Negotiations - Thru Bluff and Bluster	• Value Engineering	• Cost Targeting
	• Part Proliferation	• Standardization	• Reduced Part No. and Model Base	• Business Concentration
	• Random Part No. System	• Preferred Parts List	• Description Data Base	• Group Technology
Cycle Time (J.I.T.)	• Large Safety Stocks	• M.R.P. and Over-Reliance on Computer	• Focused Factories	• Schedule Linearity
	• Long Lead Time	• Push System	• Pull System	• Indirect Labor Productivity
	• Poor and Yo-yo Forecasts	• Long Production Runs • Long Set-up Time • Process Flow	• Small Lot Sizes, Short Set-up and Use • Product Flow	• Near-Instant Customer Delivery

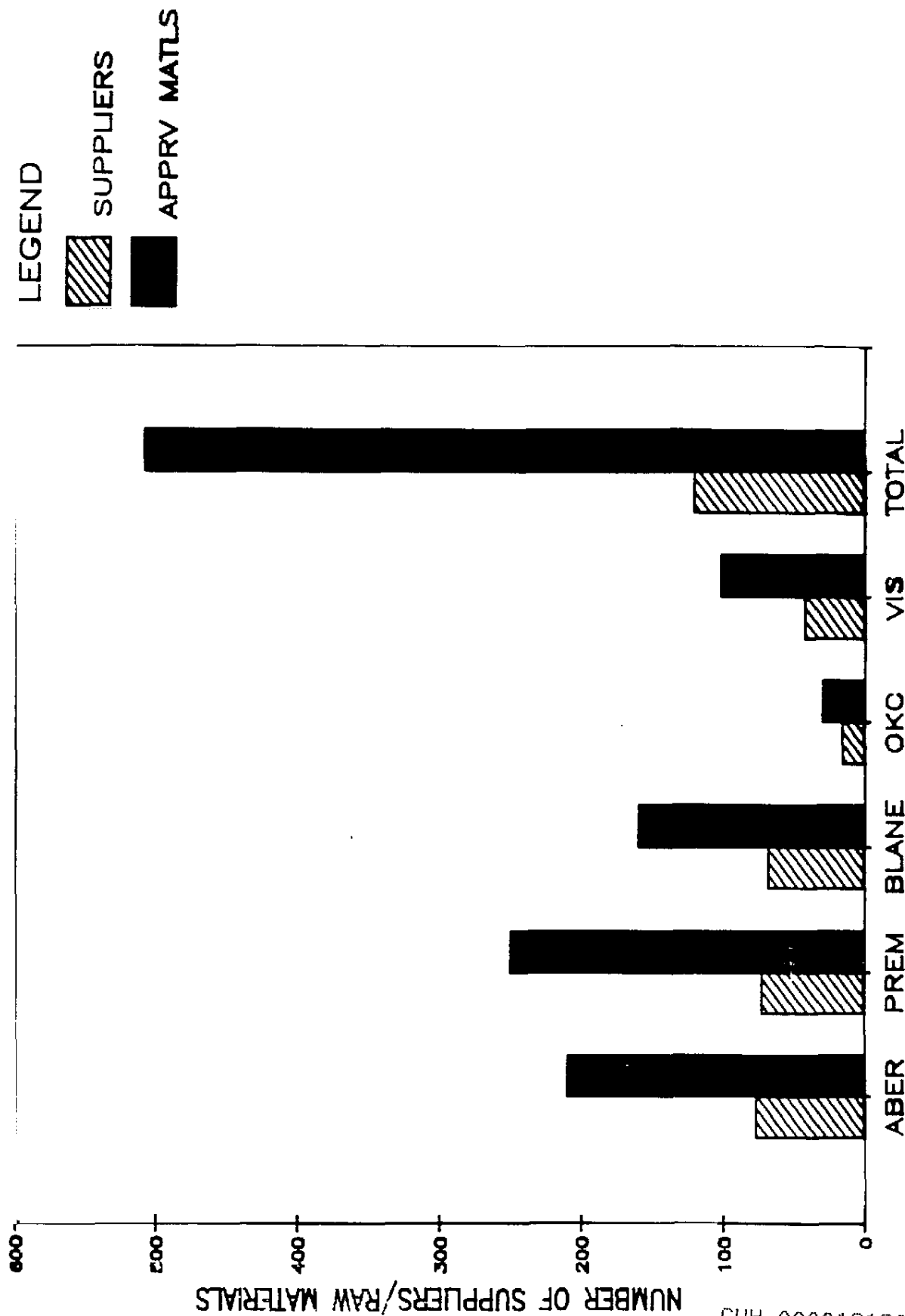


PROBLEM IN ESTABLISHING

MEANINGFUL SUPPLIER PARTNERSHIPS ...

TOO MANY RAW MATERIALS AND SUPPLIERS !!!

POLYMERS APPROVED RAW MATERIALS



WHAT DO WE WANT TO ACHEIVE???

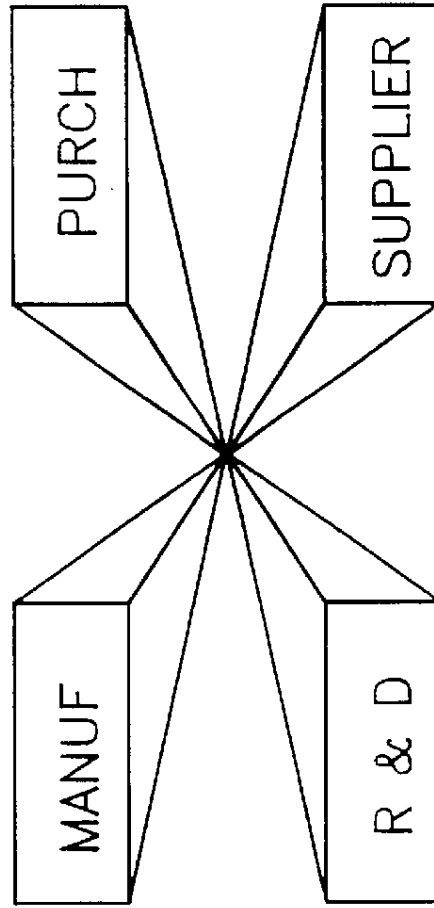
- o IMPROVE RAW MATERIAL QUALITY
- o REDUCE RAW MATERIAL VARIATION
- o REDUCE COSTS THROUGH LONGER-TERM
HIGHER-VOLUME CONTRACTS
- o REDUCE INVENTORY COSTS
- o REDUCE COSTS IN PURCHASING ADMINISTRATION
- o ENHANCE R & D/TECHNICAL ASSISTANCE
AVAILABLE TO VISTA

FUTURE WORK

1. Establish Commodity Teams
2. Continue Supplier Surveys
3. Develop Intimate Supply Partnerships
4. Develop an Overall Raw Material Supply Process

ESTABLISH COMMODITY TEAMS

- o Organize within different Business Areas
- o Make **appropriate** personnel available

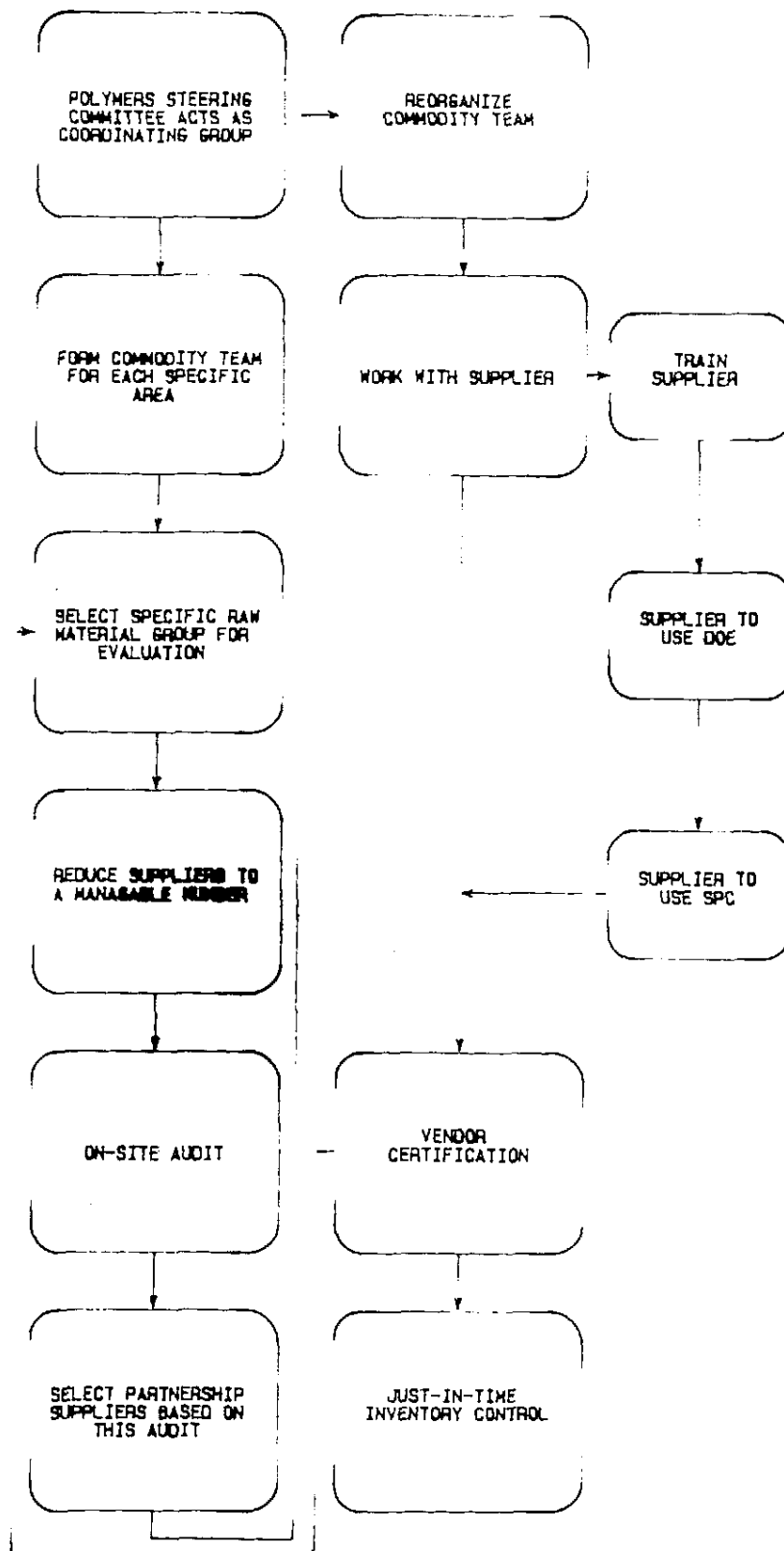


SUPPLIER EVALUATION AND SELECTION

"... the knowledge required to select suppliers that will be good candidates for a long-term relationship **directed** toward continuous improvement does not and will not reside exclusively within the purchasing function."

Ohio Quality and Productivity
Forum Roundtable Conference

SUPPLIER EVALUATION
PROPOSED PROGRAM



1988 SURVEYS COMPLETED

SURFACTANTS

ALUMINUM POWDER	1
CHLORINE	3
BENZENE	1
KEROSENE	1
SPECIALTY	1

POLYMERS

LEAD STABILIZERS	2
TIN STABILIZERS	3
PLASTICIZERS	1
CLAYS	2
INITIATORS	2

MRO/PACKAGING/SPECIALTY

PACKAGING	9
DRUMS	2
WATER TREATMENT	2
COMPUTER PAPER	2
MRO SUPPLIES	2
	===
	34

1989 SURVEYS PLANNED

SURFACTANTS

ALUMINUM POWDER	2
KEROSENE	3
BENZENE	2
ETHYLENE OXIDE	1
ALUMINUM CHLORIDE	3

POLYMERS

ANTIMONY TRIOXIDE	2
INITIATORS	1
CALCIUM STEARATE	2
PHTHALIC ANHYDRIDE	3
IMPACT MODIFIERS	2

MRO/PACKAGING/SPECIALTY

MRO	9
PACKAGING	3
DISTRIBUTORS	4
SPECIALTY	3
	==
	48

DEVELOP INTIMATE SUPPLY PARTNERSHIPS

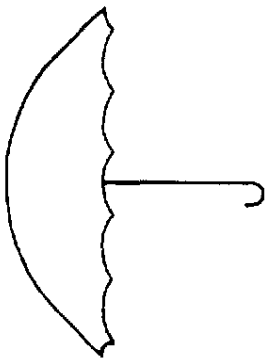
1989 GOALS:

One Commodity Team establishes a detailed partnership with one supplier.

Two other Commodity Teams are initiated.

1990 and BEYOND:

Continue reducing the number of suppliers and establishing partnerships.



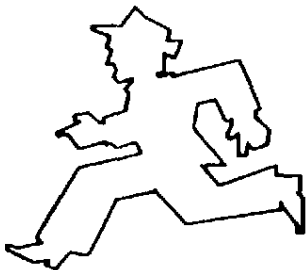
DEVELOP AN OVERALL RAW MATERIAL SUPPLY PROCESS

1. Supplier evaluation and certification process
2. Raw material approval process

1989 MANPOWER REQUIREMENTS

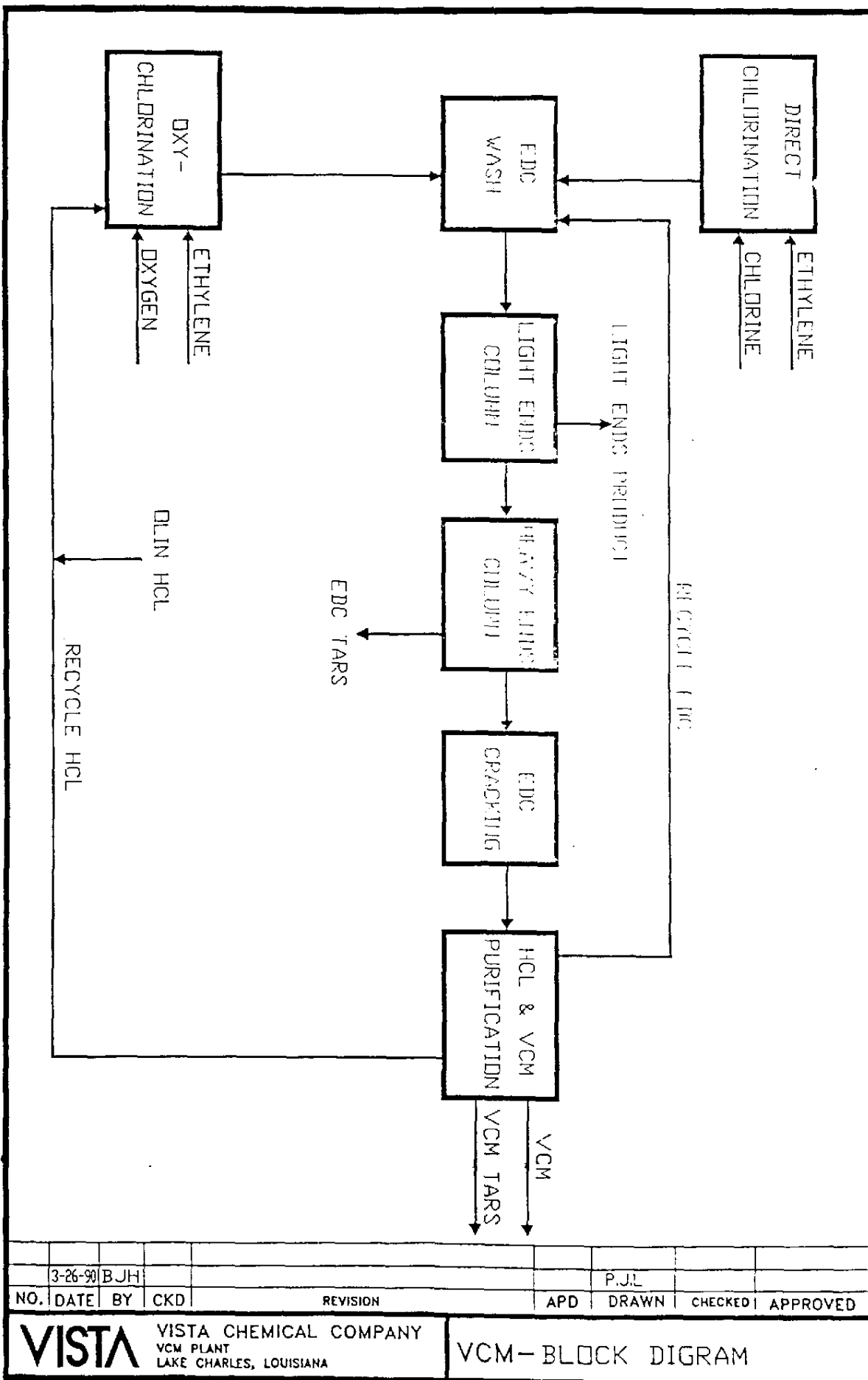
- o Purchasing 0.25–0.5 man–years
- o Manufacturing 0.25–0.5 man–years
(Excluding test–runs, etc.)
- o R & D 0.25–0.5 man–years
(Excluding any extensive lab work)

CMH 0000121A0



For Each Raw Material

- 1) Do we receive COA's?
- 2) Which parameters are important / which vcm plant parameters are affected by supplier purity - pick top two
- 3) Do we analyze?



CWH 000012141

Statistical Summary

Sphere Analyses
May Through June, 1991

<u>Component</u>	<u>LCL</u>	<u>Mean</u>	<u>UCL</u>
Alkalinity, ppm as NaOH	0.015	0.306	0.597
Iron, Nonfilterable ppm	-0.039	0.034	0.108
Water, ppm	6.0	55.1	104.3
Acetaldehyde, ppm	-0.072	0.141	0.354
Acetylene, ppm	-0.138	0.044	0.226
1,3-Butadiene, ppm	5.050	7.087	9.124
C4 Unsaturation, ppm	11.180	16.665	22.151
Methyl Chloride, ppm	26.148	45.185	64.222

Tank Car Analyses
May Through "mid" July, 1991

<u>Component</u>	<u>LCL</u>	<u>Mean</u>	<u>UCL</u>
Oxygen Before Loading, ppm	-129.8	175.2	480.3
Oxygen After Loading, ppm	-60.0	85.5	231.0

Lake Charles VCM Plant

QMP History

- 1986 Full Time Coordinator — Facilitate efforts / be a resource
 QIP Training - Salaried
 -Quality Philosophy
 -SPC
 -Problem Solving
- 1987 IPSM Training - All Employees - develop better understanding/enhance working relationships
 QIP Training - Hourly Personnel
 Operations Charting Begins
 Lab QA Team Formed - establish QA for critical areas
 -Lab Charting Begins - provide valid analytical results
 Experimental Design Training Begins
- 1988 Team Management Training Begins - Use teams as a Forum for process improvement
 For Salaried Personnel
 -Teamwork
 -Problem Solving
- 1989 Natural Team Startup - all employees involved in process
- 1990 Gainsharing - share in gains made
 Team Management Training - Maint.
 Operations Charting Revised - never made any tremendous headway
 -realized need to make it effective
 -just recently put a system in which should begin to move us in the right direction

9-28-90

To: All team members, HOA, WPB, SMS

Re: Tenth meeting of the pipefitter-welder natural team

Attendees: GLL,BTH,BB,HJA,LBT,KWV,JNF,LD,GMS(facilitator)

The meeting opened in due form at 7:30 a.m., with Garland serving as team leader. The minutes of the previous meeting were read and reviewed. The code of conduct was reviewed with no discussion brought up--none was needed. We went directly into the agenda, which was shop safety problems. Nelson noted that the breathing air system had been installed in the shop, and was something that had been needed for some time. Luke brought up that a manifold system for the oxy-acetylene torches was needed to eliminate the problem of all the "burning" rigs being scattered around in the shop. This causes a congested work area, as well as other problems. Joel pointed out that housekeeping during major outages such as turnarounds, equipment failures, etc. cause additional safety problems. Barry mentioned the traffic problem in the shop area--too many people passing through without safety equipment--(sales people, visitors, etc.) Tom brought up the storage bin on the south wall--it needs to be revised, the shop grinder needs warning signs and shields replaced, the N2 test table needs to be organized with racks, etc. being installed and labeled, and the problem with other crafts leaving junk and other materials in the pipe shop. Kenneth mentioned that people using portable grinders need to be aware of the direction of their "spark stream", as this could cause injury to others in the area. Garland said that we need a storage cabinet for combustible materials used in the shop, and set up a pm program for stationary equipment, we also have a problem with electrical cords on fans, and other equipment causing tripping hazards--we need to look at an overhead plug system. On the second round of brainstorming, Barry pointed out that contaminated materials need to be cleaned before bringing into the shop. Koon said that the roof exhaust fan needs to be repaired or replaced, and that it is too small anyway.

We then did a recap and decided we would begin working on problem solving as follows; Koon and Kenneth--check on ventilation and flammable storage problem; Joel and Larry--work on housekeeping and test table problem; Barry and Luke--traffic and electrical cord problem; Garland, Tom, and Shoe--equipment "pm" program, oxy-acet. header, and storage bin problems. The consensus was that we would begin working on these problems ASAP and have a progress report ready for the next meeting which was set for Oct. 10. Having no further business to conduct, the meeting adjourned at 8:10 am.


Luke Darter, Recording Secretary

CWH 000012144

Meeting Agenda

Meeting Subject: GIL SAN MARCOS NATIONAL TRAIL MEETING #12

Date/Time: 9-28-90 2:30 PM. To 3:30 PM.

Place: SOUTH CONFERENCE BUILDING

Attendees: J.B., K.W., R.L., P.B., T.C. (Facil) S.M.S.

Purpose/Objectives: ① IDENTIFY TOP POTENTIAL SAFETY HAZARDS
② IDENTIFY CAUSES,
④

Agenda Item	Responsibility	Time
Review of last meeting*	Leader	2 MINS
Code of Conduct	Facil./Leader	1 MINS
Problem - Solving Model**	Facil./Leader	2 MINS
Statement of Objectives	Leader	1 MINS
<u>Review the STATPROCESS FLOW</u>		
<u>DIAGRAM</u>		<u>13 MINS</u>
<u>IDENTIFY THE NUMBER OF</u>		
<u>POTENTIAL HAZARDS</u>		<u>10 MINS</u>
<u>SET A GOAL TO DECREASE</u>		
<u>POTENTIAL HAZARDS</u>		<u>5 MINS</u>
<u>IDENTIFY THE HAZARDS TO</u>		
<u>WORK ON FIRST</u>		<u>10 MINS</u>
<u>ANALYZE THE PROCESS</u>		<u>10 MINS</u>
<u>DETERMINE CAUSES</u>		<u>4 MINS</u>
Action Steps/Schedule	Leader	
Adjourn	Leader	<u>2 MINS</u>

*If not the first meeting.

**If meeting is the first of a problem solving team.

- Reminder:
1. Start/End on time.
 2. Designate a secretary.
 3. Distribute meeting notes.

Distribution: J.B., K.W., R.L., P.B., T.C., S.M.S., H.O.A., W.P.R.

CWH 000012145

Boiler Maker NATURAL TEAM Meeting #11
DATE & TIME 8-24-90 11:30 AM TO 12:30 PM.

11:30 ATTENDANCE: J.B. - S.S. - R.L. - K.W. -
P.B. - T.C.

11:35 COOR OF CONDUCT

11:40 REVIEW LAST MEETING:

11:50 REVIEW GRAPH FOR PERFORMANCE
OF JOB & SAFETY.

12:00 QUALITY OF NAOH BEING HAND
IN BARREL'S WAS DISCUSSED.

12:10 SUGGESTED REVISION TO NAOH
LOADING CHECK SHEET TO BE
ENLARGED TO SEE BETTER.

12:12 SUGGESTION TO "FLAG" OFF AREA
WHEN LOADING NAOH IN DRYER.

12:15 MOTION TO SEND OUT LETTER TO
PLANT PERSONNEL TO HONOR OUR
FLAGGING WHILE LOADING NAOH
IN DRYER'S.

12:25 NEXT MEETING TO RE-VALUE THE
NAOH CHECK SHEET

12:30 NEXT MEETING AFTER TURN-A-ROUND

CAUSTIC LOADING CHECK SHEET

DATE	9/13																		
First Aid / Injury																			
NEAR MISS	⑧																		
JSA Steps not done	2																		
JSA HAZ Situations	1.3 4.5																		
Unresolved CASES																			
SAFETY Total																			
# OF LEAKS																			
Amount of Correct CAUSTIC																			
START Time	7:09																		
STOP Time	7:55																		
TOTAL																			

- ① JSA HAZ. SIT. - Press. on Bott. Line.
- ② JSA STEPS - BLIND on Bott. not removed to Depressure
- ③ NO FACE Shield - 2EA
- ④ NOT Lowering FACE Shield - Band
- ⑤ STANDING UNDER Load - Band
- ⑥ AREA NOT FLAGGED off - People WALKING THROUGH AREA.
- ⑦ BARREL SOFT AWAY - CAUSTIC FLEW - BAD. CAUSTIC

CWH 000012147

CAUSTIC LOADING CHECK SHEET

DATE	9/19																		
First Aid / Injury																			
NEAR MISS																			
ISA Steps not done	1/																		
ISA Haz Situations	2/																		
Unresolved CASES																			
SAFETY Total																			
# OF LEAKS																			
Amount of Correct CAUSTIC																			
START Time	7:00																		
STOP Time																			
TOTAL																			

- ① S.S. VALVE NOT UNLOCKED OR OPENED
- ② STANDING UNDER LOAD
- ③ FACE SHIELDS NOT DOWN AT PROPER TIMES

①

Solutions

Causes

Potential Hazards

Job Steps

Get Permit			
Get Tools			
Roll Blinds to Blind			
Open for Manway			
Estimate Caustic			

None

None

• Caustic Burns (dripping)

• None

• None

2

Solutions

Causes

Potential Hazards

Job Steps

Bring in
Caustic with
Forklift

Unload OFF
Pallets

Bring in
Flatbed
Truck

Set up
~~potstake~~
Crane

Set Funnel
on top
of dryer

• None

• Pinned Fingers
• Mashed Toes

• None

• None

• None

Job Steps

Send up
caustic drums

Move caustic
around on
ground with
2 wheel drum
dolly

Set Funnel
Down and
Wash caustic
out of it

Close top
Manway

Roll blinds
open

Potential Hazards

Drums could fall
From air

Drum could fall
off onto someone

Drips of caustic
Flakes falling

None

caustic burns

Causes

Solutions

Job Steps

Potential Hazards

Causes

Solutions

Unload Flat
Bed Truck

Move Crane
Out of
Block

Wait For
Good leak
check

Turn in
Tools

• None

• None

• None

• None