



Document Review Cycle

REVIEW PERIOD	NEXT REVIEW	DOCUMENT OWNER
N/A	N/A	Contracts Manager

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Revision History

YYYY	MMM	DD	lssued By	Rev	Detail
2018	JAN	08	C. Taylor	C1	First issue on BMS

Revision Change Notices

Rev	Location of Changes	Brief Description of Changes

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1.0 COMPANY INFORMATION

Company Name:	
Company Address:	
Phone Number:	Fax Number:
Number of years in business:	Number of employees:
Name of EnQuest Account Representative:	Title:
This survey is completed by:	Title:
Describe the Products or Services your company pro	vides EnQuest:

2.0 QUALITY SYSTEM REGISTRATION

2.1 Does the organisation have a third party quality system registration?

Yes 🗌 No 🗌

System:	Certification No.	Expiration date:
System:	Certification No.	Expiration date:
System:	Certification No.	Expiration date:

Comments:

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3.0 HEALTH, SAFETY, ENVIRONMENT AND ASSURANCE

3.1	Does your company have a documented HSE & A Policy?	Yes 🗌	No 🗌
3.2	Has your HSE & A process been audited and approved by any recognised		
	organisation within the past 2 years?	Yes 🗌	No 🗌
3.2.1	If so, please provide the name of auditing company, date and standard		
	you have been approved to?		
3.3	Is this HSE & A policy known by all employees?	Yes 🗌	No 🗌
3.4	Is an organisational chart or other means of defining and communicating		
	authority and responsibility within the organisation available?	Yes 🗌	No 🗌
3.5	Is there a management representative for HSE & A issues?	Yes 🗌	No 🗌
3.5.1	Name: Title:		
3.6	Does a defined system exist for communicating customer requirements		
	between departments?	Yes 🗌	No 🗌
3.7	Are management reviews of the quality management system held?	Yes 🗌	No 🗌
3.8	Are management reviews record maintained?	Yes 🗌	No 🗌
3.9	When necessary, are formal corrective actions defined, implemented, verified		
	and closed out as a result of these reviews?	Yes 🗌	No 🗌
3.10	Does the organisation have an effective means to measure customer		
	satisfaction?	Yes 🗌	No 🗌
3.11	Do HSE & A personnel have the authority to stop operation for cause?	Yes 🗌	No 🗌
Commen	ts:		

3.12	Does your company have a Safety Management Interface Document with		
	EnQuest?	Yes 🗌	No 🗌
3.12.1	If so, how has the content of this document been communicated to your		
	workforce?		
3.13	Is your company's Environmental Management System certified against ISO		
	14001 Standard?	Yes 🗌	No 🗌
3.14	What are the company environmental objectives as part of the Environmental		
	Management System, and do you do anything beyond Environmental Legal		
	Compliance (e.g. the minimum requirements)?	Yes 🗌	No 🗌
3.15	If applicable, how does the company identify, inspect and maintain		
	Environmental Critical Equipment?	Yes 🗌	No 🗌
3.16	Has your company suffered any statutory notifiable environmental invident in		
	the past year? If so, please provide details.	Yes 🗌	No 🗌
Commen	ts:		

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4.0 HSE&A RESPONSIBILITY AND AUTHORITY

4.1	Does the department responsible for HSE&A report directly to the		
	Head of organisation?	Yes 🗌	No 🗌
4.2	Does the department responsible for Quality have the authority to		
	control further processing and delivery of products and services		
	until the unsatisfactory condition has been corrected?	Yes 🗌	No 🗌
Comme	ents:		

5.0 CONTRACT REVIEW

5.1	Are all customer contracts / purchase orders reviewed prior to start of		
	work?	Yes 🗌	No 🗌
5.2	Is the department responsible for HSE & A involved in this review?	Yes 🗌	No 🗌
5.3	If a problem arises which may impact the quality, quantity, or delivery		
	of the contract, is the customer notified?	Yes 🗌	No 🗌
5.4	Are operations suspended until written response is obtained from		
	customer?	Yes 🗌	No 🗌
Co	mments:		

6.0 INTERNAL AUDITS AND CONTROL OF RECORDS

6.1	Are internal audits or self-assessments performed with results		
	documented and reported to management?	Yes 🗌	No 🗌
6.2	Are records of internal audits kept?	Yes 🗌	No 🗌
6.3	Are the retention periods of the records specified?	Yes 🗌	No 🗌
6.4	Are records adequately protected against damage or accidental		
	destruction?	Yes 🗌	No 🗌
6.5	Are procedures defined to determine the responsibility for record		
	disposition post retention period?	Yes 🗌	No 🗌
Com	ments:		

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7.0 DOCUMENTS CONTROL

7.1	Are there procedures related to the control of documents?	Yes 🗌	No 🗌
7.2	Do these procedures contain facility for document review, update and		
	approval?	Yes 🗌	No 🗌
7.3	Is there a system in place for document revision identification?	Yes 🗌	No 🗌
7.4	Are documents available at the Office?	Yes 🗌	No 🗌
7.5	Is there a process in place to preclude the unintended use of obsolete		
	documents?	Yes 🗌	No 🗌
7.6	Are revision histories posted with the documents?	Yes 🗌	No 🗌
7.7	Is the distribution of documents of internal and external origin controlled?		
	(Standards, customer drawings, etc.)	Yes 🗌	No 🗌
7.8	When external documents are used in the Office, are there procedures in		
	place to ensure that the latest revision is available?	Yes 🗌	No 🗌
Con	nments:		

8.0 COMPETENCY AND TRAINING MANAGEMENT

8.1	Have minimum competency requirements been established?	Yes 🗌	No 🗌
8.2	Is training, education and work experience for personnel documented?	Yes 🗌	No 🗌
8.3	Are there formal training programs for individuals whose work has a direct		
	effect on quality?	Yes 🗌	No 🗌
8.4	Are validation records for process or service provision available?	Yes 🗌	No 🗌
8.5	Is the organisation's production equipment on a scheduled maintenance		
	program?	Yes 🗌	No 🗌
8.6	Are planned maintenance shutdowns addressed in production		
	scheduling?	Yes 🗌	No 🗌
8.7	Is maintenance and repair work conducted in accordance with		
	documented procedures?	Yes 🗌	No 🗌
8.8	Are maintenance and repair work documented and records up to date?	Yes 🗌	No 🗌
8.9	Where you supply to EnQuest personnel in positions identified as		
	minimum positions for Competency Assessments as specified in the Oil		
	and Gas UK Guidelines, please provide evidence of such assessments.		
	Have you provided evidence as an attachment?	Yes 🗌	No 🗌
Comr	nents:		

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8.10 Please also explain how you ensure that the competency and training are up to date and meet the required standard.

Please add the detailed explanation as an attachment, where necessary.

Expl	anation:	
8.11	Please also provide a copy of the competency matrix (record) for EnQuest Platforms and Modules.	r personnel deployed at all
	Have you provided the copy as an attachment?	Yes 🗌 No 🗌
Expl	anation:	

9.0 CONTRACT AND ORDER REVIEW

9.1	Is there a documented procedure in place for Purchase order and contract		
	review to ensure customer requirements are met?	Yes 🗌	No 🗌
9.2	Do the procedures include a clear means for communicating to the		
	customer any build or specification issues?	Yes 🗌	No 🗌
9.3	Are these procedures always followed?	Yes 🗌	No 🗌
9.4	Are exceptions to contract requirements resolved before acceptance of a		
	contract?	Yes 🗌	No 🗌
9.5	Are order and contract reviews documented?	Yes 🗌	No 🗌
9.6	Is there a procedure in place for communicating changes to a customer		
	order among affected departments?	Yes 🗌	No 🗌
9.7	Is there a procedure in place for communicating product changes to the		
	customer which may affect operation, obsolescence or safety?	Yes 🗌	No 🗌
Con	nments:		

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10.0 INSPECTION TESTING AND VERIFICATION

10.1	Do supplied materials and components undergo inspection, test or other		
	means of verification upon receipt?	Yes 🗌	No 🗌
10.2	Are procedures documented and implemented for incoming materials		
	inspections all the time?	Yes 🗌	No 🗌
10.3	Is the system designed to prevent the incorporation of uninspected		
	materials into equipment or processes under most circumstances?	Yes 🗌	No 🗌
10.4	Are inspection processes and test procedures in written form?	Yes 🗌	No 🗌
10.5	When required by contract, does the organisation control the		
	implementation of customer's procedures?	Yes 🗌	No 🗌
10.6	Is the inspection status of the product readily apparent?	Yes 🗌	No 🗌
10.7	Has the organisation documented and implemented final test and		
	inspection procedures?	Yes 🗌	No 🗌
10.8	Are all tests and inspections documented and records maintained,		
	including nonconforming results?	Yes 🗌	No 🗌
10.9	Is the organisation's quality management system designed to prevent the		
	final release of un-inspected equipment or processes under most		
	circumstances?	Yes 🗌	No 🗌
Com	ments:		

11.0 PROCEDURAL CONTROLS

11.1	Are the following process controls available?:	Yes 🗌	No 🗌
	Development of personnel with essential and documented training or		
	qualifications and certifications.	Yes 🗌	No 🗌
	Adequate equipment.	Yes 🗌	No 🗌
	Controlled working environment.	Yes 🗌	No 🗌
	Defined acceptable and unacceptable work standards.	Yes 🗌	No 🗌
	Issuance of and compliance with work instructions, procedures,		
	specifications and drawings.	Yes 🗌	No 🗌
	Monitoring of process compliance with procedures.	Yes 🗌	No 🗌
Com	ments:		

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12.0 CUSTOMER OWNED PROPERTY

12.1	Is there process in place to identify customer property?	Yes 🗌	No 🗌	
12.2	Are there adequate facilities available to store and protect customer			
	property?	Yes 🗌	No 🗌	
12.3	Are records maintained of reporting lost, damaged or unacceptable	Yes 🗌	No 🗌	
	materials to the customer who supplied them?			
Comments:				

13.0 CONTROL OF NON-CONFORMING EQUIPMENT

13.1	Are there documented procedures for the control of nonconforming				
	products?	Yes 🗌	No 🗌		
13.2	Do the procedures call for the identification, documentation, evaluation				
	and handling of the nonconforming product?	Yes 🗌	No 🗌		
13.3	Are these procedures implemented?	Yes 🗌	No 🗌		
13.4	Do all customers generate NCRs reviewed by the HSE & A department?	Yes 🗌	No 🗌		
13.5	Does the organisation have an effective system for communicating NCR				
	(Non Conformance Report) status with the customer?	Yes 🗌	No 🗌		
13.6	Are instances of nonconforming product documented?	Yes 🗌	No 🗌		
13.7	Are reworked items reinspected and retested?	Yes 🗌	No 🗌		
13.8	Are records of any reinspection maintained and available?	Yes 🗌	No 🗌		
13.9	Are NCR investigated and actions shared with customers?	Yes 🗌	No 🗌		
Com	Comments:				

14.0 REMEDIAL AND PREVENTIVE ACTION

14.1	Are there documented procedu	ures for investigating and implementing)	
	corrective actions to prevent re	ecurrence of non-conformance?	Yes 🗌	No 🗌
14.2	Are all non-conformances subj	ected to these procedures?	Yes 🗌	No 🗌
14.3	Are records of corrective action	ns maintained and shared?	Yes 🗌	No 🗌
14.4	Are records of preventive actio	ons available?	Yes 🗌	No 🗌
14.5	Does the organisation have an	effective means of measuring continue	al	
	improvement?		Yes 🗌	No 🗌
14.6	Are corrective action plans revi	iewed by senior management before th	neir	
	implementation?		Yes 🗌	No 🗌
14.7	Is the progress of corrective/pr	eventive action monitored?	Yes 🗌	No 🗌
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14.8	Are corrective/preventive actions reviewed for effectiveness when		
	completed?	Yes 🗌	No 🗌
14.9	Are effective corrective/preventive actions reflected in revised		
	procedures?	Yes 🗌	No 🗌
Com	ments:		

15.0 SUPPLIER MANAGEMENT

15.1	Do you have a documented process for selecting and evaluating			
	suppliers?	Yes 🗌	No 🗌	
15.2	Are supplier assessments and reviews documented and records			
	maintained?	Yes 🗌	No 🗌	
15.3	Is there a list of approved suppliers maintained and available?	Yes 🗌	No 🗌	
15.4	Are purchases only made from approved suppliers?	Yes 🗌	No 🗌	
15.5	Is supplier performance periodically re-evaluated?	Yes 🗌	No 🗌	
15.6	Do suppliers go through formal pre-qualification process before			
	placement of contracts or orders?	Yes 🗌	No 🗌	
15.7	Do you have a process for regular supplier audits and performance			
	management?	Yes 🗌	No 🗌	
Comments:				

16.0 WAREHOUSING, PACKAGING AND DELIVERY

16.1	Are there processes in place for handling, storing, packaging and		
	protecting the product?	Yes 🗌	No 🗌
16.2	Are materials handled in accordance with documented procedures?	Yes 🗌	No 🗌
16.3	Are the procedures designed to protect the integrity of the handled		
	material as well as personnel?	Yes 🗌	No 🗌
16.4	Is appropriate handling equipment available as and when needed?	Yes 🗌	No 🗌
16.5	Are storage areas and conditions appropriate to the nature of the goods		
	held in them?	Yes 🗌	No 🗌
16.6	Are inward and outward movements from warehouse areas controlled		
	and properly authorized?	Yes 🗌	No 🗌
16.7	Are components with limited shelf life audited and evaluated routinely?	Yes 🗌	No 🗌
16.8	Are records of these audits maintained and available?	Yes 🗌	No 🗌

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16.9	Are customer requirements for packaging reviewed and evaluated for	Yes 🗌	No 🗌
	adequacy?		
16.10	Does packaging, including identification, comply with all relevant		
	statutory requirements?	Yes 🗌	No 🗌
16.11	Are delivery methods evaluated to ensure that they preserve the quality		
	of the supplied items?	Yes 🗌	No 🗌
16.12	Is product quality preserved between final test and inspection and		
	acceptance by the customer?	Yes 🗌	No 🗌
Comme	ents:		

17.0 ANTI-CORROPTION AND BRIBERY

17.1	Does your company have a regularly monitored anti-corruption code of				
	conduct documenting your anti-corruption and bribery policies and				
	procedures?	Yes 🗌	No 🗌		
17.2	Is this anti-corruption code of conduct known and acted upon by all				
	employees of your company?	Yes 🗌	No 🗌		
17.3	Is this anti-corruption code of conduct publicised internally and				
	externally?	Yes 🗌	No 🗌		
17.4	Does your company carry out documented risk assessments of its				
	potential exposure to corruption and bribery?	Yes 🗌	No 🗌		
17.5	Does your company apply due diligence procedures for persons who				
	perform or will perform services for or on behalf of the organisation, to				
	mitigate identified bribery risks?	Yes 🗌	No 🗌		
Comme	nts:				

18.0 COMPLIANCE WITH MODERN SLAVERY LEGISLATION

18.1 Is your company required by section 54 of the Modern Slav				
	produce an annual statement s	setting out the steps that have been tal	ken	
	to ensure your company and s	upply chains are slavery free?	Yes 🗌	No 🗌
18.2	Does your company have a po	licy on modern slavery and human		
	trafficking? (this may form part	of your company's wider CSR policy)	Yes 🗌	No 🗌
18.3	Does your company carry out	documented risk assessments of its		
	business and supply chain to e	ensure continuous compliance with		
	relevant statutory requirements	s?	Yes 🗌	No 🗌
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Comments:_____

LIST OF DOCUMENTS REQUIRED

Copy of your HSE & A Policy	
Copy of your training, qualification and work experience for personnel	
Copy of the competency matrix (record) for personnel deployed at all EnQuest	
Platforms and Modules	
Copy of your documented procedures for investigating, planning and	
implementing corrective actions to prevent recurrence of non-conformance	
Copy of your incident investigation process	
Copy of your incident investigation form	
Copy of your anti-corruption code of conduct	
Copy of your statement complying with section 54 of the Modern Slavery Act	
2015, if applicable	

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CONCLUSION

This area is for EnQuest internal use onl	у.	
Reviewed by:	Date:	
Reviewed by:	Date:	
Reviewed by:	Date:	
Status	Completion Date	Attachments
APPROVED		Yes 🗌 No 🗌
CONDITIONALLY APPROVED		Yes 🗌 No 🗌
CORRECTIVE ACTION		Yes 🗌 No 🗌
REQUESTED		Yes 🗌 No 🗌
NOT-APPROVED		

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