CONTRACT DATA REQUIREMENTS LIST

(1 Data Item)

Form Approved
OMB No. 0704-0188

The public reporting burden for this collection of information is estimated to average 110 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden, to the Department of Defense, Executive Services and Communications Directorate (0704-0188). Respondents should be aware that notwithstanding any other provision of law, no person shall be subject to any penalty for failing to comply with a collection of information if it does not display a currently valid OMB control number. **Please do not return your form to the above organization. Send completed form to the Government Issuing Contracting Officer for the Contract/PR No. listed in Block E.**

A. CONTRACT LINE ITEM NO.	B. EXHIBIT	C. CATEGORY: TDP _____ TM _____ OTHER _____

D. SYSTEM/ITEM	E. CONTRACT/PR NO.	F. CONTRACTOR
	HT9402-10-C-0002	

1. DATA ITEM NO.	2. TITLE OF DATA ITEM	3. SUBTITLE	17. PRICE GROUP
N/A	Q100 Evolving Practices Report		NSP

4. AUTHORITY *(Data Acquisition Document No.)*	5. CONTRACT REFERENCE	6. REQUIRING OFFICE	18. ESTIMATED TOTAL PRICE
N/A	Policy Manual (TPM), Chap 1 Sect 2.1	TMA-OCMO/MB&RS/TRO/Prog Ops	0.00

7. DD 250 REQ	9. DIST STATEMENT REQUIRED	10. FREQUENCY	12. DATE OF FIRST SUBMISSION
NO	B	QTRLY	See Block 16
8. APP CODE		**11. AS OF DATE**	**13. DATE OF SUBSEQUENT SUBMISSION**
N/A		N/A	See Block 16

16. REMARKS

Block 12 - No later than 15 calendar days following the end of the first calendar quarter in which health care delivery occurs.
Block 13 - No later than 15 calendar days following the end of each calendar quarter.

The contractor shall keep abreast of new and evolving technologies and bring to TMA's attention via submission of a quarterly report, any device, medical treatment, or procedure they believe has moved from unproven to proven.

For each device, medical treatment, or procedure reported as having moved from unproven to proven, the contractor must submit actual copies of the articles, studies, technology assessment(s), etc., which substantiates their reported conclusion. A citation or reference page is not sufficient, and only reliable evidence as defined in 32 CFR 199.2 is acceptable.

TMA will review the supporting documentation provided, apply the standards and procedures in TRICARE regulation and policy, and make the final determination concerning the proven status of the devices, medical treatments, and/or procedures included in each quarterly report submitted by the contractor.

14. DISTRIBUTION

a. ADDRESSEE	b. COPIES Draft	Final Reg	Final Repro
Submit through the			
TMA E-commerce			
Extranet			
15. TOTAL →	0	0	0

G. PREPARED BY	H. DATE	I. APPROVED BY	J. DATE
Carol L. McCourt		Laura L. Sells	

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