

**REPORTING AND PROCESSING MEDICAL MATERIAL COMPLAINTS/
QUALITY IMPROVEMENT REPORT**

1. DATE
2. NO.

3. TO	4. FROM
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5. COMPLAINT

a. TYPE <input type="checkbox"/> QUALITY <input type="checkbox"/> NEW ITEM <input type="checkbox"/> SIMILAR ITEM	b. FOR DOD USE <input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III
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6. MANUFACTURER STOCK NO.	7. ITEM DESCRIPTION
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8. MANUFACTURER

a. NAME	9. NAME OF CONTRACTOR (If other than the manufacturer)
b. ADDRESS	10. CONTRACT NO. OR PURCHASE ORDER NO.
	11. LOT NO.

12. CONTROL NO.	13. MANUFACTURER'S SERIAL NO.	14. MODEL NO.
15. DATE MANUFACTURED	16. DATE PACKED	17. EXPIRATION DATE
18. SOURCE (Direct or distributor)	19. QUANTITY ON HAND	20. QUANTITY SUSPENDED

COMPLETE ITEMS 20-25 BELOW FOR DOD TYPE I COMPLAINTS ONLY

21. TOTAL NO. PATIENTS INVOLVED	22. TOTAL NO. REACTIONS	23. SEVERE OR UNUSUAL REACTIONS
24. REACTIONS REQUIRING HOSPITALIZATION	25. LENGTH OF HOSPITALIZATION	26. VACCINE
		INITIAL
		BOOSTER

27. CAUSE OF COMPLAINT (Explanation of unsatisfactory condition, deficiency, or description of reaction. Complete for ALL complaints.)

28. INITIATOR

a. NAME (For Type I MC/DC/NC)	b. DSN TELEPHONE NUMBER		c. COMMERCIAL TELEPHONE NUMBER		
	AREA CODE	PHONE NUMBER	AREA CODE	PHONE NUMBER	EXT.

29. SUPPLY OFFICER

a. SIGNATURE	b. DATE	d. DSN TELEPHONE NUMBER		e. COMMERCIAL TELEPHONE NUMBER		
c. NAME		AREA CODE	PHONE NUMBER	AREA CODE	PHONE NUMBER	EXT.

30. REPORTING AND PROCESSING MEDICAL MATERIAL COMPLAINTS/QUALITY IMPROVEMENT REPORT (Continued)

a. RECOMMENDATIONS AND/OR ADDITIONAL REMARKS

b. ACTION TAKEN

31. ACTION OFFICER

a. NAME	b. TITLE	c. ORGANIZATION	d. DATE