A Watson Pharma, Inc.

Call Center Operations
Call Center Operational Procedure

PROCEDURE:	Suspicious Orders of Controlled Drugs		
WRITTEN BY:	Judy Callahan	DATE:	May 3, 2004
Call Center Policy Number:	OPDCC 507509-03.00	Policy Effective Date	May 3, 2004
REVISION WRITTEN BY:		REVISION DATE:	
Supersede CC Policy #	CC 1007.00	CC Policy Date:	09/03
cSOP Reference #	cSOP 11-004	cSOP Ref. Date:	May 3, 2004

 Purpose: To assure distribution of controlled drugs is monitored for excessive use by an individual location using the DEA number as the identifier.

 Scope: This procedure applies to all controlled drugs distributed by Watson Laboratories, Inc. and its Subsidiaries.

## DOCUMENT REFERENCES:

Document Number	Document Title		
CTMAN-80-041-CC-OPR	Order Processing		
CTMAN-80-041-CC-OPS	Order Processing Supervisor		

### ATTACHMENTS:

Document Number	Document Title	
N/A	N/A	

## DEFINITIONS:

	DEA	Drug Enforcement Agency	
• A	SOMS	Suspicious Order Management System	

#### III. Procedure:

#### 1.0 Process for Suspicious Orders of controlled drugs

Responsibility	Action	
General	1.1	The SAP system compiles a past history of controlled substance drug product orders by each customer to establish a normal order size and order frequency. This is accomplished through the normal Sales Order process see CTMAN- 80-041-CC-OPR, Order Processing, for details on this process.
Call Center Management/ Controlled Substance Compliance Management	1.2	The SOMS Multiplier Table is determined by Call Center Management and the Controlled Substance Compliance Department.

CONFIDENTIAL - SUBJECT TO PROTECTIVE ORDER

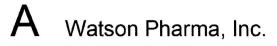


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# Call Center Operations Call Center Operational Procedure

PROCEDURE:	Suspicious Orders of Con	spicious Orders of Controlled Drugs		
WRITTEN BY:	Judy Callahan	DATE:	May 3, 2004	
Call Center Policy Number:	OPDCC 507506-03.00	Policy Effective Date	May 3, 2004	

Order Processing Representative	1.3	<ul><li>1.2.1 See CTMAN 80-045-CC-LEM, License Entry and Management for description of system functionality with regards to the establishment of SOM levels, as well as how to access this information.</li><li>The list is to be updated by written request to the customer approximately 30 days prior to the expiration date of the current registrations.</li></ul>
License Administrator	1.4	The license administrator will review the SOMS report, and then contact the customer to confirm the quantity ordered and verify the reason for a larger or more frequent order.
	1.5	Once this SOMS report is confirmed and verified by the Customer, the SOMS report is signed and marked with a reason code by the license administrator and submitted to the Supervisor or Management for review and signature.
	1.6	The license administrator will be responsible to ensure that pending sales orders on hold due to suspicious order (SOMS) violation are investigated.
	1.7	The license administrator will release pending orders due to SOMS violations by canceling the order, or reducing the quantity, per customer requirements.
	1.8	If the SOMS violation cannot be resolved by canceling the order or reducing the quantity, the license administrator will escalate the suspicious order to the next level.
	1.9	Determine if the order does or does not classify as suspicious.
	1.10	If a valid reason (based on objective criteria) does not exist, the order will be deemed as a suspicious order and will not be filled. Report suspicious issue to Control Substance Compliance Department.
Controlled Substance Compliance Department	1.11	The control Substance Compliance Department will determine the next level of Communication.
Order Processing Representative	1.12	File a copy of the SOMS Report, along with the customer purchase order, in the suspicious order record file.