OMB—SUPPORTING STATEMENT

Federal Ceiling Price Retail Pharmacy Program

A. Justification

1. <u>Need for Information</u>

Pursuant to the terms of a contract awarded by DoD, a commercial pharmacy benefits manager (PBM) will provide a retail pharmacy network for the DoD TRICARE Management Activity. The PBM will issue payment with Government funds for prescriptions dispensed by retail network pharmacies to TRICARE beneficiaries. DoD will provide manufacturers with itemized data on covered drugs purchased through TRICARE retail network pharmacies in order to obtain appropriate refunds on covered drugs delivered to TRICARE beneficiaries.

2. <u>How, by Whom, and for What Purpose Information Will be Used</u>

DoD will provide manufacturers with itemized data on covered drugs purchased through TRICARE retail network pharmacies in order to obtain appropriate refunds on covered drugs delivered to TRICARE beneficiaries.

3. <u>Does information collection involve the use of Information Technology</u>

As indicated above DoD will provide manufacturers with itemized data on covered drugs purchased through TRICARE retail network pharmacies. The drug manufacturers will validate the refund based on the difference between a benchmark price, consisting of either the manufacturer's actual sales price to the wholesaler or retail pharmacy chain when known and auditable or non-FAMP (non-Federal average manufacturer price) and the Federal Ceiling Price (FCP).

The manufacturers currently use IT to download and decrypt their data and going forward will utilize the web-based system to obtain their data and submit their disputes.

4. Efforts to Avoid Duplication

There are no existing data which can be used for these purposes.

5. <u>Small Business Impact</u>

None.

6. <u>Consequences of Not Collecting Information</u>

If the proposed data collection is not approved, DoD not be able to obtain appropriate refunds on covered drugs delivered to TRICARE beneficiaries.

7. Special Circumstances

There are no special circumstances involved in this data collection effort.

8. Applicability to 5 CFR 1320.8(d)

The draft notice was published in the *Federal Register* on February 25, 2009 (74 FR 8512-8513). No comments were received.

The Pharmaceutical Operations Directorate contacted several manufacturers to verify how long it took them to complete the tasks of downloading, decrypting, reviewing the data, disputing claims and submitting payments.

9. Remuneration to Respondents

No payments or gifts will be provided to respondents.

10. **Confidentiality**

Data will be used solely for the purpose of obtaining appropriate refunds on covered drugs

delivered to TRICARE beneficiaries and for no other purpose.

11. Questions of a Sensitive Nature

There are no questions of a sensitive nature included in the collection.

12. Hour Burden Estimate

There are an approximate total of 250 drug manufacturers responding to this collection. There

will be four responses per year per respondent with an estimated 8 hours of preparation time per response

for a total of 8,000 hours.

AFFECTED PUBLIC: Approximately 250 Drug Manufacturers who have entered into a master

agreement with the VA.

ANNUAL BURDEN HOURS: 8,000

NUMBER OF RESPONDENTS: 250

RESPONSES PER RESPONDENT: 4

AVERAGE BURDEN PER RESPONSE: 8 hours

FREQUENCY: Quarterly

8,000 hours x \$60.09 (representative hourly professional salary including benefits) =

\$480,720.00

13. Capital, Start-up, and Maintenance Costs

Respondents will be asked to maintain records. No additional equipment purchases will be made

to support data collection processes or record keeping, thus no incremental cost above the cost of the

collection of the information will be incurred.

3

14. Annualized Cost to Federal Government

The total cost to the government for collecting this information \$84,060.

15. Changes from OMB Form 83-I

This is a reinstatement of a previously approved collection; therefore, the change in burden is 8,000.

16. Outside Publication

The data collected will be used for internal DoD use only and there are no plans for outside publication of results.

17. Expiration Date

The expiration date of the OMB approval will be displayed on the information collection.

18. <u>Certification Statement</u>

The proposed data collection does not involve any exceptions to the certification statement identified in line 19 of OMB Form 83-I. As required, an agency disclosure statement will be prominently displayed on the information collection.