PUSHCHASE ORDER
THE TEXAS A&M UNIVERSITY SYSTEM
HEALTH SCIENCE CENTER
200 Technology Way, Suite 2079, College Station, Texas 77845-3424; Phone 979-436-9219, FAX 979-436-0074

P700037

VENDOR GUARANTEES MERCHANDISE DELIVERED ON THIS ORDER WILL MEET OR EXCEED SPECIFICATIONS IN THE RFP INVITATION.

ALL TERMS AND CONDITIONS SET FORTH IN THE RFP INVITATION BECOME A PART OF THIS ORDER.

TEXAS A&M HEALTH SCIENCE CTR
DEPT OF NEUROSCIENCE AND EXPERIMENTAL THERAPEUTICS
MEDICAL RESEARCH EDU BLDG
8447 STATE HIGHWAY 47 STE 1005
BRYAN TX 77807-3260

SHIP TO:

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VENDOR QUOTE: 827564 R-2

TOTAL 94,000.00

FOB: DESTINATION FRT INCLUDED

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PURCHASING AGENT FOR
THE TEXAS A&M UNIVERSITY SYSTEM HEALTH SCIENCE CENTER

The State of Texas is exempt from all Federal Excise Taxes

STATE AND CITY SALES TAX EXEMPTIONS CERTIFICATE: The undersigned claims an exemption from taxes under Texas Tax Code, Section 151.366(d), for purchase of tangible personal property described in this numbered order, purchased from contractor and/or shipper listed above, as this property is being secured for the exclusive use of the State of Texas. The Terms and Conditions of the State of Texas shall prevail.
**PURCHASE ORDER**

THE TEXAS A&M UNIVERSITY SYSTEM

HEALTH SCIENCE CENTER

FILE

200 Technology Way, Suite 2079, College Station, Texas 77845-3424; Phone 979-436-9219, FAX 979-436-0074

Order Date: 01/09/2017

Page 02

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**ITEM**

**DESCRIPTION**

Purchase made by an Institution of Higher Education, Section 51.9335 Education Code.

**CC**  **FY**  **ACCOUNT NO.**  **DEPT.**
23  2017  123002-00011-5453  2010
23  2017  462674-00001-5453  2010

**DOCUMENT DATE:** 01/09/2017

**DEPT. CONTACT:** TAMIE SEYDLER

**PHONE NO.:**

**SOLE SOURCE REASON:**

NO OTHER INSTITUTIONS

**PCC CD:** 9

**TYPE FUND:** S  **TYPE ORDER:** HIED

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THE TEXAS A&M UNIVERSITY SYSTEM HEALTH SCIENCE CENTER
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THE TEXAS A&M UNIVERSITY SYSTEM HEALTH SCIENCE CENTER
**PURCHASE ORDER**

**THE TEXAS A&M UNIVERSITY SYSTEM HEALTH SCIENCE CENTER**

200 Technology Way, Suite 2079, College Station, Texas 77845-3424; Phone 979-436-9219, FAX 979-436-0074

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**VENDOR**

P700037

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**INVOICE (IN DUPLICATE) TO AGENCY BELOW**

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**Purchasing Agent for THE TEXAS A&M UNIVERSITY SYSTEM HEALTH SCIENCE CENTER**

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THE TEXAS A&M UNIVERSITY SYSTEM HEALTH SCIENCE CENTER

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Signature: [Signature]

Date: [Date]
Finally,

I have your PO, however I can't find a vendor email address.
Please forward the attached to the vendor.

Thanks,

Susan
This form is to be used to aid departmental staff in relating information necessary in the process of requisitions on a sole source and/or proprietary basis. Your cooperation in answering the questions listed below will assist the purchaser in handling your order expeditiously. Please complete the form and forward to HUB and Procurement Services. If more space is required, feel free to attach additional pages. NOTE: For your convenience, this is a fill-in form. Adobe Reader required.

1. Description of item (if commodity: make, model no., etc.; if service: detail of type of service):
   Seizure and PK Studies in the Soman Model in Rats

2. Name of known source for item: Manufacturer? □ Yes □ No
   MRIGlobal, a non-profit research service institution, Kansas City, MO

3. What feature or functions are unique (proprietary) to this item?
   The proposed services involves toxic chemical agents that are exclusively available from MRIGlobal, which has license from US Department of Defense.

4. Briefly explain how the unique features or functions are essential to the purpose for which the item is needed.
   The proposed soman model is essential to screen new medications for this chemical neurotoxicity. This study is an integral part of the FY04 studies in the NIH grant awarded to Dr. Reddy (TAMHSC).

5. List any source other than the known source that manufactures or supplies similar items or items with similar functions.
   None from any other institutions.
Seizure and PK Studies in the Soman Model in Rats

Letter Proposal

Texas A&M University Health Science Center

D. Samba Reddy, Ph.D., RPh, FAAPS, FAAAS
Professor
Neuroscience and Experimental Therapeutics
College of Medicine
Texas A&M University Health Science Center
MREB Bldg., Rm No. 2008, 8447 State Highway 47
Bryan, Texas 77807-3260

MRIGlobal Proposal No.
827564 R-2

December 21, 2016

National Security • Global Health • Energy
Missouri • Colorado • Florida • Maryland • Virginia • Kansas • Washington, D.C.
425 Volker Blvd., Kansas City, Missouri 64110-2241 • Phone: 816-753-7600, Fax: 816-753-8420 • www.mriglobal.org
December 21, 2016

D. Samba Reddy, Ph.D., RPh, FAAPS, FAAAS
Professor
Neuroscience and Experimental Therapeutics
College of Medicine
Texas A&M University Health Science Center
MREB Bldg., Room No. 2008, 8447 State Highway 47
Bryan, TX 77807-3260

Subject: MRIGlobal Proposal No. 827564 R-2, “Seizure and PK Studies in the Soman Model in Rats”

Dear Dr. Reddy:

MRIGlobal is pleased to submit this revised proposal to the Texas A&M University Health Science Center entitled, “Seizure and PK Studies in the Soman Model in Rats.” Included in this proposal are the Technical Approach; Programmatic Assumptions; Deliverables; Performance Schedule; Price and Payment; and Project Authorization. The modification reduces the animals and time points on the PK studies (Task 1 and 2) from 45 animals to 40 and from 9 time points to 8. Dr. Reddy has also agree to send up to two of his staff to help with activities on Task 3 and 4.

Technical Approach

The details provided in this proposal represent MRIGlobal’s understanding of the work and the estimated cost associated with performance of the Statement of Work (Appendix A). MRIGlobal and Texas A&M University Health Science Center agree that this proposal will be modified only if significant changes that affect cost are requested and that these changes will be addressed via formal notification by authorized individuals for both parties.

Programmatic Assumptions:

- MRIGlobal will procure non-EEG rats (from Taconic) for these studies, and will get IACUC approvals.
- TAMU will ship EEG telemetry rats to MRIGlobal for these studies.
- MRIGlobal will have a 2 week quarantine/acclimation period and will send 1-2 serum samples to CRL for serology as the animals are coming from a University rather than a vendor.
- A technician from the Reddy lab will be on site for 2 days prior to the first study to train MRIGlobal technical staff on the video/telemetry equipment.
• Dr. Reddy and/or his technician will be on site for the first 24 hr of the first study; however, they are not permitted in the RDS laboratory, but will be on site to watch video feed from the lab and close by for technical input if trouble shooting of video/telemetry equipment is necessary.
• Shipping of all telemetry/video/equipment and animals, including fixed tissues at the conclusion of the study, will be paid for by Dr. Reddy.
• MRIGlobal is not responsible for any disposable parts of the equipment and if necessary, these parts will be replaced by Dr. Reddy.
• All animals will be euthanized at 24 hr after challenge with soman and perfused; brain will be harvested and shipped back to the Reddy lab for histology at Reddy lab.
• A new soman sample will be prepared after one year if subsequent animal studies take place after the first preparation is made; subsequent (annual) preparations of soman would be added by contract modification if this occurs as this was not included in this ROM or SOW.
• This study will not be conducted under GLP guidelines, however MRIGlobal SOP’s and GLP documentation procedures will be followed.
• The study protocol or study amendments will be authored, reviewed and signed by the Sponsor prior to execution of all studies. The study protocol will be approved by the Sponsor and MRIGlobal’s IACUC prior to ordering animals for delivery.

**Deliverables**
MRIGlobal will provide a letter report and tissues for each task stated herein.

**Performance Schedule**
MRIGlobal is proposing to perform this work with a period of performance beginning December 21, 2106 through April 30, 2017. The proposed 5-month period of performance includes letter report, archiving, and administrative close out.

**Price and Payment**
The total fixed price to complete the effort proposed herein is outlined in Table 1 on the following page.

MRIGlobal will invoice each study/milestone as priced in Table 1 at completion. The total invoice is due within thirty (30) days of the invoice date. MRIGlobal reserves the right to discontinue the work in the event payments become delinquent. MRIGlobal will provide Texas A&M University Health Science Center with a written notice within five (5) days before discontinuing work because of nonpayment of invoices.

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Project Authorization

If the proposed scope of work, price and period of performance described above are acceptable, MRIGlobal would anticipate receipt of a subcontract from Texas A&M University Health Science Center authorizing the work. MRIGlobal reserves the right to negotiate the terms and conditions along with the Statement of Work of any such subcontract.

If you have technical questions regarding this proposal please contact Mr. Ed Sistrunk at (816) 326-5650, or via email at esistrunk@mriglobal.org. Questions on contractual issues may be addressed to the undersigned at (816) 326-5012, or via email at rcole@mriglobal.org.

Sincerely,

MRIGlobal

[Signature]

Ed Sistrunk, MSM, PMP
Associate Division Director
Medical Countermeasures Division

Approved for:

MRIGLOBAL

[Signature]

Rosilyn Cole
Senior Contracts Administrator
Appendix A.
Statement of Work
Title: Seizure and PK Studies in the Soman Model in Rats Task 1:

Background
Organophosphate nerve agents such as soman (GD) and VX produce lethal neurotoxic effects, such as hypersecretion, tremors, convulsions, respiratory distress and death. Nerve agent toxicity is treated with 2-PAM, atropine sulfate, and diazepam, which suppress convulsions and seizures when given early. The longer these seizures progress, the more resistant they become to diazepam, resulting in greater neuropathology. Reddy lab team is investigating new treatments for control of nerve agent toxicity, especially convulsive seizures and brain damage. The team would like to test neurosteroids, which are potent GABA-A receptor modulators of with anticonvulsant properties, against the nerve agent-induced seizures and neuropathology in rats. The team would also like to study PK of neurosteroids in rats exposed to soman.

Objectives
The main objective of this study is to determine the effect of the neurosteroid analogs against soman-induced seizures and neurotoxicity in rats (non-GLP studies). Another objective is to determine the PK of neurosteroid analogs in soman-exposed rats. The proposed soman model will be conducted at MRIGlobal facility under a collaboration/contractual arrangement.

Experimental Design
1. Male Sprague-Dawley rats (250-400 g) will be implanted with a pair of EEG recording electrodes in the brain in the Reddy lab at TAMHSC in Bryan, Texas.

2. After a post-surgical recovery period of at least 2 weeks, these rats will be shipped to the MRIGlobal facility for soman exposure studies. MRIGlobal will care for these rats. After successful clearance of quarantine health check, rats will be utilized for studies.

   NOTE: MRIGlobal will procure non-EEG rats (from Taconic) for these studies, and will get IACUC amendments to their AUPs covering these compounds.

3. On the day of testing, rats will be hooked up to the video-EEG system for a 30-min baseline recording.

   NOTE: The video-EEG system will be supplied by Reddy lab (shipped to MRIGlobal a few days in advance), and training given to MRIGlobal technicians for operating the system. Reddy lab technician will be available on site for assistance in running video-EEG during the entire study period.

4. Rats will be given the cholinesterase agent HI-6 (125 mg/kg, i.p.) 30-min before soman exposure.

   NOTE: HI-6 (125 mg/ml) vial will be supplied by Reddy lab.

5. Rats will be exposed to soman (154 μg/kg, 1.4xLD₃₀) by a single subcutaneous injection (Ref. Figueiredo et al., J Pharmacol Exp Ther 2011,336:303-312).

6. MRIGlobal will prepare soman (154 μg/ml) solution in ice cold saline for injection. It is injected to rats at a volume of 0.1 ml per 100gm body weight or in a volume not to exceed 0.3 ml per animal.
7. Within 1 min after **soman** exposure, rats will receive atropine methyl nitrate (2 mg/kg, i.m.).
   
   **NOTE:** AMN (2 mg/ml) vial will be supplied by Reddy lab.

8. Test drugs Ganaxalone will be given **40 min after** exposure to **soman**. The study groups will be organized into 2 or 3 sessions as outlined below:

   **Study Session I (January-2017) – PK study (Sample Collection only)**

   **Task 1: Day-1:**

   Group I: **Neurosteroid GX-21** (10 mg/kg, i.m. injection) at 40-min post-GD, 5 rats/group (5x8 time points, N=40)

   **NOTE:** A **20 ml vial of neurosteroid Gx-21** (10 mg/ml) injection solution will be supplied by Reddy lab. A single shot is to be given 40-min post-GD (40 rats x 0.3 ml = 12.0 ml needed for this study)

   **Sample Collection:** See PK Protocol below.

   **Task 2: Day-2:**

   Group II: **Neurosteroid GX-Lysine** (10 mg/kg, i.m. injection) at 40-min post-GD, 5 rats/group (5x8 time points, N=40)

   **NOTE:** A **20 ml vial of neurosteroid Gx-Lysine** (10 mg/ml) injection solution will be supplied by Reddy lab. A single shot is to be given 40-min post-GD (40 rats x 0.3 ml = 12.0 ml needed for this study)

   **Sample Collection:** See PK Protocol below.

   **PK Study Protocol:**

   - Forty (40) min post-GD injection, each rat will receive an injection I.M. (intramuscular) of test **neurosteroid** at a dose of 10 mg/kg, divided over two thigh muscle sites (each thigh) in a volume of 0.2-0.3 mL (not to exceed 0.3 ml per site).
   
   - Clinical observations including onset of seizure activity will be recorded throughout study period (from GD exposure till sample collection for all groups).
   
   - After the administration of neurosteroid, four (5) rats per time-point will be anesthetized with an IP injection of ketamine and xylazine then exsanguinated.
   
   - Ketamine and xylazine: Anesthetics administered by IP injection and at target doses of ≤ 100 to 200 mg/kg (Ketamine) and ≤ 20 mg/kg (xylazine), respectively. **NOTE:** Rats experiencing GD-induced seizures activity may need more Ketamine dose for inducing anesthesia.
   
   - Blood sample (3-5 ml) is collected via cardiac puncture or vena cava or by external carotid artery puncture. Samples will be collected at the following time-points: **0, 15, 30, 45, 60-min, 2h, 4h and 8h** after Neurosteroid injection (total 8 groups). Time 0 is defined as time of Neurosteroid administration.
   
   - Brain tissue harvested at the same (above) time-points. **NOTE:** Allow enough time between groups to anesthetize, collect blood and brain tissue.
- In two groups (4h and 8h groups), rats will be video monitored from the time of GD injection until anesthetized (at 4h or 8 h after ganaxolone dosing) for blood and brain tissue collection. Dr. Reddy lab will provide the video system (as was done in earlier study).

- Blood will be collected via cardiac puncture or vena cava or external carotid artery puncture under ketamine/xylazine anesthesia.

  - Following blood collection into purple top EDTA tubes, the blood will be processed to plasma, then transferred into a minimum of two labeled tubes and stored at -80°C ± 10°C. **NOTE:** Collect the plasma in two separate (set A and B) tubes (0.5 to 1 ml each tube) and store at -80°C in plasma storage tubes.

  - Ship one set of plasma samples (set A) to SRI for analysis.

  - Store the second set (set B) at MRIGlobal at -80°C.

- The brain following harvest will be divided into a right and left half (midline), each half weighed separately, and then placed into either tubes or seal-safe labeled bags and placed onto dry ice.

  - At the end of tissue harvest, brain samples will be transferred to and stored at -80°C. **NOTE:** Weight of each both tissues (set A & set B) taken immediately after harvesting and recorded on the sheets (before cooling the tissue in dry ice).

  - Ship one set of brain samples (set A) to SRI for analysis.

  - Store the second set (set B) at MRIGlobal at -80°C.

**Study Session II (February-2017) – EEG study (regular study)**

**Task 3: Week-1 (February-2017) – Neurosteroid Analogs study**

**Day-1 & 2:**

Group I: **Neurosteroid GX-21** (10 mg/kg, i.m.) at 40-min post exposure, N= 15 rats (7 EEG + 8 non-EEG)

**NOTE:** A 30 ml vial of neurosteroid Gx-21 (5 mg/ml) injection solution will be supplied by Reddy lab. Two shots are to be given 40-min post-GD; additional boosters shots at 30-min & 1 or 2 h after the first shot (15 rats x 0.3 ml x 2 shots = 9 ml per dose; 3 doses =30 ml needed).

Group II: **Neurosteroid GX-Phosphate** (20 mg/kg, i.m.) at 40-min post exposure, N= 15 rats (7 EEG + 8 non-EEG)

**NOTE:** A 30 ml vial of neurosteroid Gx-Phosphate (10 mg/ml) injection solution will be supplied by Reddy lab. Two shots are to be given 40-min post-GD; additional boosters shots at 30-min & 1 or 2 h after the first shot (15 rats x 0.3 ml x 2 shots = 9 ml per dose; 3 doses =30 ml needed).

**Day-3 & 4:**

Group III: **Neurosteroid GX-Lysine** (20 mg/kg, i.m.) at 40-min post exposure, N= 15 rats (7 EEG + 8 non-EEG)
NOTE: A 30 ml vial of neurosteroid Gx-Lysine (10 mg/ml) injection solution will be supplied by Reddy lab. Two shots are to be given 40-min post-GD; additional boosters shots at 30-min & 1 or 2 h after the first shot (15 rats x 0.3 ml x 2 shots = 9 ml per dose; 3 doses = 30 ml needed).

Group IV: Neurosteroid Gx-Valine (20 mg/kg, i.m.) at 40-min post exposure, N= 15 rats (7 EEG + 8 non-EEG)

NOTE: A 30 ml vial of neurosteroid Gx-Valine (10 mg/ml) injection solution will be supplied by Reddy lab. Two shots are to be given 40-min post-GD; additional boosters shots at 30-min & 1 or 2 h after the first shot (15 rats x 0.3 ml x 2 shots = 9 ml per dose; 3 doses = 30 ml needed).

Task 4: Week-2 (February-2017) – Combination study

Day-1 & 2:

Group I: Neurosteroid Gx-Phosphate+Midazolam (2 mg/kg, i.m.) at 40-min post-GD, N= 15 rats (7 EEG + 8 non-EEG)

NOTE: A 30 ml vial of neurosteroid Gx-Phosphate (10 mg/ml) injection solution will be supplied by Reddy lab. Two shots are to be given 40-min post-GD; additional boosters shots at 30-min & 1 or 2 h after the first shot (15 rats x 0.3 ml x 2 shots = 9 ml per dose; 3 doses = 30 ml needed).

NOTE: Midazolam injection (2 mg/ml) will be prepared by MRIGlobal.

Group II: Neurosteroid Gx-Lysine+Midazolam (2 mg/kg, i.m.) at 40-min post-GD, N= 15 rats (7 EEG + 8 non-EEG)

NOTE: A 30 ml vial of neurosteroid Gx-Lysine (10 mg/ml) injection solution will be supplied by Reddy lab. Two shots are to be given 40-min post-GD; additional boosters shots at 30-min & 1 or 2 h after the first shot (15 rats x 0.3 ml x 2 shots = 9 ml per dose; 3 doses = 30 ml needed).

NOTE: Midazolam injection (2 mg/ml) will be prepared by MRIGlobal.

NOTE: Test drugs solutions will be provided by Reddy lab in vials ready for administration (0.1 ml per 100 g body weight of animals). Test drugs should be stored in the refrigerator.

NOTE: Midazolam (5 mg/ml) injection vials should be procured by MRIGlobal from veterinary suppliers (Controlled substance).

9. Rats are monitored continuously by Video-EEG recording system for 24 h after soman exposure.

NOTE: In each group, there will be 7 rats hooked to the video-EEG recording system another 8 rats video recording only (non-EEG animals), total of 15 rats in each group.

NOTE: Rats can be observed in their home cages, which can be fitted with swivels for free movement of animal during studies.

NOTE: Video-EEG data will be recorded in the PC provided by Reddy lab.
10. Rats will be perfused transcardially with 0.4% paraformaldehyde at 24 hours after soman exposure. A few animals (N=5 to 6), non-EEG rats, from each group are perfused at 24 h post exposure.


11. Brains will be delicately isolated, processed as per the perfusion protocol, and samples stored in 0.4% paraformaldehyde for histological studies.

   **Note:** Brain samples will be shipped to Reddy lab after completion of fixation procedures.

12. All surviving rats will be transferred to animal facility, cared intensely with fluids and moistened food for 3-5 days post exposure. Upon recovery, animals will be shipped to Reddy lab, preferably 1 or 2 weeks after the study.

   **Note:** Health reports, safety and related information are needed prior to shipping the rats to Reddy lab.

**Specific Deliverables:**

**Session 1: PK study**

Samples will be shipped to SRI on dry ice (with prior notification) the week following PK study execution.

A copy of the study file will be shipped to Dr. Reddy upon completion of this program.

**Session 2: EEG study**

Brain samples will be shipped by overnight FedEx to Reddy lab.

Surviving animals will be shipped to Reddy lab, preferably 1 week after the study.

**Reporting Requirements. None**

Research technician from Reddy lab will be at the site for the video-EEG data collection.
Purchasing

DEBARRED VENDOR LIST

The following vendors shown below are debarred from doing business with the State of Texas, effective from the date of debarment for the length of time indicated. Whether they are listed below or not, the debarred vendors include the vendors' successors in interest as defined in Rule §20.102(b)(4).

<table>
<thead>
<tr>
<th>Vendor ID Number</th>
<th>Vendor Name/Address</th>
<th>Date of Debarment</th>
<th>Length of Debarment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1562456928900</td>
<td>Smith Housewares and Restaurant Supplies 500 Erie Blvd. Syracuse, NY 13202</td>
<td>November 12, 2014</td>
<td>5 Years</td>
</tr>
<tr>
<td>1743261315000</td>
<td>Walker's Electric Company 1520 Park St Beaumont TX 77701 Also: Walkers Electric Company Calvin G. Walker Stacy Walker</td>
<td>August 28, 2012</td>
<td>5 Years</td>
</tr>
</tbody>
</table>
Federal Exclusion

Agencies and co-op members may wish to check the list of vendors excluded from doing business on the federal level. The System for Award Management, or SAM®, can be used as a resource for purchasing entities.

According to Statewide Procurement Division rules, other debarment activities from other entities may be considered as possible indicators of vendor responsibility.

Vendor Information on Payments

The Search State Payments Issued application provides vendors with payment details.

They can also sign up in the application for Advance Payment Notification.

Texas Government Code §2155.077®

VISUAL COMPLIANCE RESTRICTED PARTY SCREENING

Search criteria: MRIGlobal  (Exact match)
[Export, Sanctions, GSA, Police, PEP and International data groups]

Date of search: Monday, January 9, 2017
Time of search: 10:05 AM EDT
Report created by: SUSAN WARREN, TEXAS A&M - HEALTH SCIENCE CENTER

NO MATCHING RECORDS FOUND

AUTHORITIES:

- Department of Commerce Denied Persons [BIS]
- Department of Commerce Entity List [BIS]
- Department of Commerce "Unverified" List [BIS]
- Department of State Arms Export Control Act Debarred Parties [DDTC]
- Department of State Munitions Export Control Orders [DDTC]
- Department of State Nonproliferation Orders
- Department of State Iran Sanctions (ISA and TRA)
- WMD Trade Control Designations [OFAC]
- Department of State Designated Terrorist Organizations
- Department of State Terrorist Exclusion List
- Palestinian Legislative Council List [OFAC]
- Federal Register General Orders
- Specially Designated Nationals and Blocked Persons [OFAC]
- Foreign Sanctions Evasion List (FSE-IR) [OFAC]
- Sectoral Sanctions Identifications List (UKRAINE-E013662) [OFAC]
- Persons Identified as Blocked Solely Pursuant to Executive Order 13599 [OFAC]
- United Nations Consolidated List
- GSA Parties Excluded from Federal Procurement Programs [SAM/EPLS]
- GSA Parties Excluded from Federal Nonprocurement Programs [SAM/EPLS]
- GSA Parties Excluded from Federal Reciprocal Programs [SAM/EPLS]
- Air Force Special Investigations - Top Ten Fugitives
- Alcohol, Tobacco, Firearms and Explosives Most Wanted
- FBI Ten Most Wanted Fugitives
- FBI Most Wanted Terrorists
- FBI Kidnappings and Missing Persons
- FBI Seeking Information
- FBI Wanted Fugitives
- Food and Drug Administration - Clinical Investigators
- Food and Drug Administration - Debarment List
- Food and Drug Administration - Disqualified and Restricted
- Homeland Security Investigations Most Wanted
- Naval Criminal Investigative Service - Wanted Fugitives
- U.S. Immigration and Customs Enforcement Most Wanted
- U.S. Drug Enforcement - Major International Fugitives
- U.S. Marshals Service - Major Fugitive Cases
- U.S. Marshals Service - Top 15 Most Wanted
- Office of Research Integrity PHS Administrative Actions
- U.S. Postal Inspection Service - Most Wanted
- U.S. Secret Service Most Wanted
- OIG Entities Excluded from Federal Health and Medicare Programs
- CIA Chiefs of State and Cabinet Members of Foreign Governments [Politically Exposed Persons]
- Japan Foreign End-Users of Concern
- Kingdom of Saudi Arabia Wanted Militants
- CPSEP Listed Entities
- Australia Foreign Affairs Consolidated List
- European Union Consolidated List
- Interpol Recently Wanted
- HM Treasury Consolidated List [England]
- Canadian Economic Sanctions
- Canadian Border Services Agency Wanted List
- RCMP Wanted Fugitives
- FinCEN (USA PATRIOT Act) Section 311 - Special Measures
- World Bank Listing of Ineligible Firms
- OSFI Consolidated List - Entities
- OSFI Consolidated List - Individuals
- OSFI Warning List