DEPARTMENT OF VETERANS AFFAIRS
Federal Supply Schedule Service
Authorized Federal Supply Schedule Price List

On-line access to contract ordering information, terms and conditions, up-to-date pricing, and the option to create an electronic delivery order are available through GSA Advantage!, a menu-driven database system.

The INTERNET address GSA Advantage! is: GSAAdvantage.gov

Schedule Title: FSC GROUP 65, PART II, SECTION A COMMODITY: MEDICAL EQUIPMENT AND SUPPLIES
FSC CLASS PRODUCT CODE: 6515 – Medical and Surgical Instruments, Equipment, and Supplies

Contract number: 36F79719D0049

For more information on ordering from Federal Supply Schedules click on the FSS Schedules button at fss.gsa.gov

Contract period: December 15, 2018 to December 14, 2023

Current through mod P00004, effective August 15, 2020

Government Marketing & Procurement, LLC
13350 Ranch Rd 12
Wimberly, TX 78676-5304

P: 703-349-2990
F: 703-995-0321
W: http://www.gmpgov.com/

Business size: Small business
Service-Disabled Veteran Owned Small business.
CUSTOMER INFORMATION:
1a. Table of awarded special item number with appropriate cross-reference to item
descriptions and awarded prices:

Special Item Number A-20C (Implants, Surgical - (c) Other)
Special Item Number A-21 D (Other Disposable Contamination Containers)
Special Item Number A-46 (Secondary Oxygen Equipment)
Special Item Number A-8A (Disposable Surgical Hand Instruments - Includes instrument accessories)
Special Item Number A-92 (Medication and Supply Packaging and Dispensing Equipment)
Special Item Number A-94 (Introduction of New Products under 65 II A, Medical Equipment & Supplies)

1b. Identification of the lowest priced model number and lowest unit price for that model for each special
item number awarded in the contract:
A-20 C Axolotl Biologix, Inc’s items ABFC05 and ABFA05 for $349.75 each.
A-21 D DisposeRx, Inc’s item NDC # 64584-0000-40 for $1,718.59.
A-46 Aireon Therapeutics, Inc’s item 9031101E for $4,666.40
A-8A Kleiner Device Labs item 10101-S for $1,432.16.
A-92 TruMed Systems, Inc’s item ReOrder for $627.14
A-94 Koelis, Inc item KIT-4011 for $954.27

1c. Hourly rates: Not applicable.

2. Maximum order: $100,000 for A-20 C, A-21 D, A-8 A and A-94 and $50,000 for A-46 and $450,000 or (1)
one system for A-92.

3. Minimum order: None.

4. Geographic coverage (delivery area): 48 contiguous States, to include the District of Columbia. Point of
Exportation (POE) to Alaska, Hawaii, and Puerto Rico.

5. Point of production: Hays County.

6. Discount from list prices or statement of net price: All discounts are listed in the price list.

7. Quantity discounts: None.


9a. Notification that Government purchase cards are accepted at or below the micro-purchase threshold:
Yes. Acceptance up to, equal to, and above the micro-purchase threshold. No limitations.

9b. Notification whether Government purchase cards are accepted or not accepted above the micro-purchase
threshold: Yes. Acceptance up to, equal to, and above the micro-purchase threshold. No limitations.
10. Foreign items: Not applicable.

11a. Time of delivery: 30 Days after receipt of order (ARO).

11b. Expedited Delivery: As mutually agreed. Ordering facilities are responsible for the difference between standard and expedited delivery.

11c. Overnight and 2-day delivery: As mutually agreed. Ordering facilities are responsible for the difference between standard and expedited delivery.

11d. Urgent Requirements: As mutually agreed. Ordering facilities are responsible for the difference between standard and expedited delivery.

12. F.O.B. point: Destination 48 contiguous States, to include the District of Columbia.
Point of Exportation (POE) to Alaska, Hawaii, and Puerto Rico.

13a. Ordering address: 13350 Ranch Rd 12, Wimberly, TX 78676-5304.

13b. Ordering procedures: For supplies and services, the ordering procedures, information on Blanket Purchase Agreements (BPA’s) are found in Federal Acquisition Regulation (FAR) 8.405-3.

14. Payment address: 13350 Ranch Rd 12, Wimberly, TX 78676-5304.

15. Warranty provision:

The below Commercial product warranties as a supplement to, not a replacement for FAR clause 52.212-4
(o) Warranty and 52.212-4 (p) Limitation of Liability.

Government Marketing & Procurement LLC accepts the Government Warranty Clauses 52.212-4 (o)
(TAILORED), and 52.212-4 (p) (TAILORED):

Warranty: The Contractor warrants and implies that the items delivered hereunder are merchantable and fit for use for the particular purpose described in this contract. In the event that the terms of the contractor’s standard commercial warranty conflict with the warranty terms contained in this clause, the terms of this clause will govern this contract, unless some other resolutions are specified in the award document.

Limitation of liability: Except as otherwise provided by an express warranty, the contractor will not be liable to the Government in a breach of warranty action for consequential damages resulting from any defect or deficiencies in accepted items. In the event that the terms of the contractor’s standard commercial warranty/limitation of liability clause(s) place greater limits on the contractor’s liability than do the terms contained in this clause, the terms of this clause will govern the contract.

GMP will honor the manufacturer’s (Axolotl) Commercial Warranty Policy.
Limited Warranty. Axolotl Biologix® warrants for a period of one year after shipment that each non-amnion based product is free from material defects in material and workmanship and has a shelf-life of at least six months from the date of shipment to Customer.

In addition, Axolotl Biologix® warrants for a period of one year after shipment that each amnion based product has been collected, processed and stored in compliance with all applicable laws and standards, including the AATB Standards and Title 21, Code of Federal Regulations 1271, Human Cells, Tissues, and Cellular and Tissue-Based Products and has a shelf-life of at least two years from the date of shipment to Customer.

DISCLAIMER OF WARRANTIES: EXCEPT AS EXPRESSLY PROVIDED IN ABOVE, SUPPLIER MAKES NO WARRANTIES OR CONDITIONS, EXPRESS, STATUTORY, IMPLIED, OR OTHERWISE, AND SUPPLIER SPECIFICALLY DISCLAIMS THE IMPLIED WARRANTIES AND CONDITIONS OF MERCHANTABILITY AND FITNESS FOR A PARTICULAR PURPOSE. NOTWITHSTANDING THE FOREGOING, AXOLOTL BIOLOGIX® DOES NOT EXCLUDE LIABILITY TO THE EXTENT THAT SUCH LIABILITY MAY NOT BE EXCLUDED OR LIMITED BY LAW.

Return Goods Policy:

Returns – 30 days to return an item. No item may be returned without a Return Merchandise Authorization (RMA) Number from our customer service department.

Non-defective items can only be returned if they were purchased directly from GMP and must be returned in unused condition and in the unopened, original packaging – due to the nature of our products, we cannot authorize the return of a product that has been opened and/or removed from its original packaging. Retail, commercial and government customers must request an RMA within 10 days of receiving order for a full refund of the purchase price.

Reason for Returns: Any Customer may return a Product to Axolotl Biologix® for 100% credit, at the Customer's option, if the Customer determines in good faith that any of the following conditions are met:

Axolotl Biologix® shipped the Product in error;

The Product is damaged before it is accepted by the Customer;

The Product packaging or crating is damaged before it is accepted by the Customer;

The Product does not materially perform to performance specifications provided by Axolotl Biologix®;

The Product does not meet industry quality standards related to performance specifications and data submissions required by the FDA or FDA approval of the Product;
The Product is outdated or expired when delivered to the Customer; or Axolotl Biologix® gives prior written approval, which must not be unreasonably withheld.

Manner of Return: Customer must contact Axolotl Biologix® to receive an RMA number and where possible, Customer must return a Product in its original packaging or crating.

Aireon Therapeutics, Inc. Return Policy
Because of the nature of our business Aireon Therapeutics Inc. does not accept returns.
If there is a problem with your order, please contact Aireon Therapeutics us using our ‘Contacts’ page (https://aireontherapeutics.net/contactus/) so we may correct any issues.

Restocking Fees: Axolotl Biologix® must not charge any restocking fee and Axolotl Biologix® must pay all return shipping costs unless the Customer ordered the Product in error.

Kleiner Device Labs Warranty
All Kleiner Device Labs’ products are warranted for product quality, and any defective products will be replaced, as long as the company’s Warranty Claim Protocol is followed. The cost of the replacement product and shipping will be paid by Kleiner Device Labs.

PRODUCT USE DISCLAIMER OF LIABILITY
KDL products are designed for use by, and only by, technically competent, trained and licensed medical professionals. Use of KDL products is at the sole discretion of a qualified treating clinician, making their own professional decisions about which surgical tools and devices to use and in what manner to use them. KDL does not dispense individual patient medical advice and recommends that surgeons be trained in the use of any particular product before using it in surgery. Damages, injuries or death caused by clinician choices of products, surgical techniques or errors are solely the responsibility of the clinician. KDL specifically disclaims any liability for clinical use, miss-use, accident, abuse, improper handling, or use for other than intended purpose. KDL’s KG 1 products are for single use only; any warranty is voided by re-use of that product. Any non-factory modifications to KDL products invalidate the warranty and immediately terminate any and all liability by KDL for the product or damages caused by its use.

WARRANTY CLAIM PROTOCOL
If you experience a product failure that you believe is a product quality, manufacturing, or sterile packaging defect, please do the following. 1) Set aside any questionable KDL product and open a new one. 2) If you believe the patient was injured by the device’s defect or failure, please call the company as soon as is practical after the procedure to report and discuss the problem first-hand. Please call 650-720-4766 x801. We may ask you to follow a different procedure if an injury is involved. 3) Only after the patient procedure has been completed, please take several photographs of the defective or questionable item, and if possible, close-up photos of the specific defect. 4) Document in writing how and at what point in the procedure the product failed. 5) Email all
photos and written description to our Quality Department at quality@kleinerlabs.com. Include your
contact information, including your phone number, in your email. 6) Dispose of the defective product as you
would normally for used, disposable instruments at your location post-procedure. Do not return used or
defective products to the company. 7) We will examine the evidence provided and will provide a replacement
product or component, as appropriate to the failure. Replacement products will be shipped to Customer at
KDL’s expense and without handling charge.

TruMed Systems Inc. Warranty, Maintenance and Privacy Policy
Warranty and Maintenance: Warranty and maintenance are provided in accordance with the terms of the
respective Platinum, Gold and Silver Service packages offered. The customer must order a Service Package with
the purchase of the AccuVax Hardware Kiosk.

Privacy Notice:
This privacy notice discloses the privacy practices for (www.trumedsystems.com). This privacy notice applies
solely to information collected by this website. It will notify you of the following: What personally
identifiable information is collected from you through the website, how it is used and with whom it may be
shared. What choices are available to you regarding the use of your data. The security procedures in place to
protect the misuse of your information. How you can correct any inaccuracies in the information.
Information Collection, Use, and Sharing We are the sole owners of the information collected on this site. We
only have access to/collect information that you voluntarily give us via email or other direct contact from you.
We will not sell or rent this information to anyone. We will use your information to respond to you, regarding
the reason you contacted us. We will not share your information with any third party outside of our organization,
other than as necessary to fulfill your request, e.g. to ship an order. Unless you ask us not to, we may contact you
via email in the future to tell you about specials, new products or services, or changes to this privacy policy.
Your Access to and Control Over Information You may opt out of any future contacts from us at any time. You
can do the following at any time by contacting us via the email address or phone number given on our website:
See what data we have about you, if any. Change/correct any data we have about you. Have us delete any data
we have about you. Express any concern you have about our use of your data. Security We take precautions to
protect your information. When you submit sensitive information via the website, your information is protected
both online and offline. While we use encryption to protect sensitive information transmitted online, we also
protect your information offline. The computers/servers in which we store personally identifiable information are
kept in a secure environment. If you feel that we are not abiding by this privacy policy, you should contact
us immediately via telephone at (844)TRUMED-1 or via email support@trumedsystems.com.

16. Export packing charges: Not applicable.

17. Terms and conditions of Government purchase card acceptance (any thresholds above the
micro-purchase level): None.

18. Terms and conditions of rental, maintenance, and repair: Not applicable.

19. Terms and conditions of installation: Not applicable.
20. Terms and conditions of repair parts indicating date of parts price lists and any discounts from list prices: Not applicable.

20a. Terms and conditions for any other services: Not applicable.

21. List of service and distribution points: Not applicable.

22. List of participating dealers: Not applicable.

23. Preventive maintenance: Not applicable.

24a. Special attributes such as environmental attributes: Not applicable.

24b. Section 508 compliance: The EIT standards can be found at: www.Section508.gov/

25. Data Universal Number System (DUNS) number: 123922788.

26. Notification regarding registration in System for Award Management (SAM) database: Yes.
Axolotl Cryo™ Product and Benefits:

Axolotl Cryo™ is a cryopreserved liquid allograft derived from the placental components of the amnion to advance soft tissue repair, replacement, and reconstruction. It is classified as minimally manipulated under FDA regulation 21 CFR Part 1271 and section 361 of the PHS.

The two primary cell lines which reside in the amnion are human mesenchymal stromal cells (hMSC) and human amnion epithelial cells (hAEC). Both of these cells are considered to be pluripotent stem cells. Axolotl Cryo™ contains growth factors and cytokines such as epidermal growth factor (EGF), vascular endothelial growth factor (VEGF), and transforming growth factor – beta (TGF-β), Interleukin-10 (IL-10).

These proteins are essential for fetal growth and development and express significant therapeutic benefits when used as a treatment to stimulate repair and regeneration. Axolotl Cryo™ is immune-privileged, anti-inflammatory, anti-fibrotic, pro-vascular, and cytoprotective because of the cells being sourced from the amnion. These secreted factors also signal endogenous progenitor cells to promote regeneration and repair of damaged or degenerated tissue.

Relevant conditions:

Axolotl Cryo™ has potential clinical benefits in a variety of applications and has been used to treat injuries such as:

**Orthopedics/Podiatry:** Tendinitis, Bursitis, Plantar Fasciitis, Ruptured Achilles Tendon, Osteo-Chondral Defects, Labral Tears Shoulder/Hip, Flexor Tendon Repair and Osteoarthritis.

**Pain Management:** Hip Abductor/Adductor Tears, Knee Injections, MCL/LCL Tears, Rotator Cuff Lesions, Epicondylitis (Tennis Elbow), Hamstring Strains/Tears, Chronic Non-Healing Wounds and Ankle Sprains.

**Quality Assurance:** The donor tissue is recovered and processed aseptically, in accordance with all FDA guidelines and quality assurance standards in a controlled environment. Axolotl Cryo™ allograft tissue products have been subjected to microbiological studies at recovery and final packaging.

Axolotl Cryo™ Product details:

- Human allograft under FDA regulation 21 CFR Part 1271 and section 361 of the PHS.
- Immune privileged with anti-inflammatory and anti-bacterial properties.
- A rich source of growth factors, proteins, cytokines, hyaluronic acid, and collagen scaffolds.
- Contains extracellular matrix components for cellular attachment and proliferation.
- Cryopreserved for an extended shelf life.
- Ease of use.

**Allograft Membrane Patch™ (aka Axolotl Graft™) Product and Benefits:**

Axolotl Graft™ is a dehydrated human amnion membrane allograft (dhAM) derived from the amniotic components of the placenta to advance soft tissue repair and reconstruction. It is classified as minimally manipulated under FDA regulation 21 CFR Part 1271 and section 361 of the PHS.

Axolotl Biologix uses our proprietary BioSym™ process to manufacture Axolotl Graft™. The amniotic components used in Axolotl Graft™ creates a natural 3-D extracellular matrix scaffold for cellular attachment and creates an environment to promote cell migration and proliferation.
Amniotic tissues are reported to contain cytokines and growth factors which stimulate protein and collagen synthesis, collagenase activity, and chemotaxis of fibroblasts and smooth muscle cells.

Axolotl Graft™ functions as a bacteriostatic agent, which can inhibit the growth of bacteria at the wound site and reduce the rate of infection in chronic wounds.

Axolotl Graft™ is immune-privileged, it lacks specific surface antigens, which makes it suitable for many different clinical applications.

Relevant conditions:

Research has demonstrated amniotic membrane products similar to Axolotl Graft™ have clinical benefits such as: Pain Reduction, Faster Healing, Anti-bacterial, Wound Adherence, Neovascularization, Immune Privilege, Less Scarring and Wound Barrier.

Quality Control: The donor tissue is recovered and processed aseptically, in accordance with all FDA guidelines and quality assurance standards in a controlled environment. Axolotl Graft™ allograft tissue products have been subjected to microbiological studies at recovery.

Axolotl Ambient™ Product and Benefits:

Axolotl Ambient™ is an ambient temperature (25±7°C, 77±13°F) stored liquid allograft derived from the amniotic components of the placenta to advance soft tissue repair and reconstruction. It is classified as minimally manipulated under FDA regulation 21 CFR Part 1271 and section 361 of the PHS.

The two primary cell lines which reside in the amnion are human mesenchymal stromal cells (hMSC) and human amnion epithelial cells (hAEC). Both of these cells are considered to be pluripotent stem cells.1 Axolotl Ambient™ contains growth factors and cytokines such as epidermal growth factor (EGF), vascular endothelial growth factor (VEGF), and transforming growth factor – beta (TGF-β), Interleukin-10 (IL-10).

These proteins are essential for fetal growth and development and express significant therapeutic benefits when used as a treatment to stimulate repair and regeneration. Axolotl Ambient™ is immune-privileged, anti-inflammatory, anti-fibrotic, pro-vascular, and cytoprotective because of the cells being sourced from the amnion. endogenous progenitor cells to promote regeneration and repair of damaged or degenerated tissue.

Relevant conditions:

Axolotl Ambient™ has potential clinical benefits in a variety of applications and has been used to treat injuries such as:

**Orthopedics/Podiatry:** Tendinitis, Bursitis, Plantar Fasciitis, Ruptured Achilles Tendon, Osteo-Chondral Defects, Labral Tears Shoulder/Hip, Flexor Tendon Repair and Osteoarthritis.

**Pain Management:** Hip Abductor/Adductor Tears, Knee Injections, MCL/LCL Tears, Rotator Cuff Lesions, Epicondylitis (Tennis Elbow), Hamstring Strains/Tears, Chronic Non-Healing Wounds and Ankle Sprains.

Quality Control: The donor tissue is recovered and processed aseptically, in accordance with all FDA guidelines and quality assurance standards in a controlled environment. Axolotl Ambient™ allograft tissue products have been subjected to microbiological studies at recovery and final packaging.
**INSTALLATION**

a. GMP will coordinate with purchasing office or any site preparations needed prior to product delivery and subsequent installation.
b. Except for preparation of the location where the system will be installed, the system cannot be installed by the customer or a contractor employed by the customer. Only certified product technicians are authorized to install the system.
c. Throughout the life of the system the purchasing office will notify GMP in writing if the place where the system is installed changes.

**WARRANTY**

a. GMP will honor manufacturer's (Koelis, Inc.) commercial warranty policy and warrants and implies that the items delivered hereunder are merchantable and fit for use for the particular purpose described in this contract.
b. GMP warrants the System and the ultrasound probes against all manufacturing defects for twelve (12) months from the date of installation on Customer's site.
c. Customer will inform GMP within 24 hours of any malfunction or defective operation, regardless of the cause.
d. In the scope of the warranty, GMP will make all necessary efforts to repair System defects and/or breakdowns within thirty (30) days of the claims being considered. If the system is unable to be repaired within 30 days GMP will coordinate delivery of replacement equipment while the System is down for repair.
e. The warranty does not cover failures related to general wear and tear, failure to follow the instruction manuals, external accidents, defective upkeep, or modifications or repairs performed by an unauthorized third party.
f. Customer's transporting or moving the System out of the installation site does not fall within the scope of normal use of the System.
g. Any repairs that are made and any parts changed during the warranty period are guaranteed until the initial warranty period expires.
<table>
<thead>
<tr>
<th>SIN</th>
<th>Part #</th>
<th>Mfg</th>
<th>Description</th>
<th>GSA</th>
<th>Warranty</th>
</tr>
</thead>
<tbody>
<tr>
<td>A-46</td>
<td>9031101C</td>
<td>Aireon Therapeutics, Inc.</td>
<td>CPAP Hydration Fluid. Pallet of Ultra-pure water specifically designed for CPAP usage. Pallet contains 96 cases, each case contains (24)-12 ounce bottles of CPAP Hydration Fluid. Using CPAC Hydration Fluid prevents growth of microorganisms (bacteria and fungus) which reduces respiratory health hazard, and increases equipment life which reduces CPAP lifecycle costs. Price includes UPS Ground shipping for customers in the Central Time Zone of the US</td>
<td>$4,910.25</td>
<td>1 Year</td>
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<td>A-46</td>
<td>9031101E</td>
<td>Aireon Therapeutics, Inc.</td>
<td>CPAP Hydration Fluid. Pallet of Ultra-pure water specifically designed for CPAP usage. Pallet contains 96 cases, each case contains (24)-12 ounce bottles of CPAP Hydration Fluid. Using CPAC Hydration Fluid prevents growth of microorganisms (bacteria and fungus) which reduces respiratory health hazard, and increases equipment life which reduces CPAP lifecycle costs. Price includes UPS Ground shipping for customers in the Eastern Time Zone of the US</td>
<td>$4,666.40</td>
<td>1 Year</td>
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<tr>
<td>A-46</td>
<td>9031101M</td>
<td>Aireon Therapeutics, Inc.</td>
<td>CPAP Hydration Fluid. Pallet of Ultra-pure water specifically designed for CPAP usage. Pallet contains 96 cases, each case contains (24)-12 ounce bottles of CPAP Hydration Fluid. Using CPAC Hydration Fluid prevents growth of microorganisms (bacteria and fungus) which reduces respiratory health hazard, and increases equipment life which reduces CPAP lifecycle costs. Price includes UPS Ground shipping for customers in the Mountain Time Zone of the US</td>
<td>$5,347.44</td>
<td>1 Year</td>
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<tr>
<td>A-46</td>
<td>9031101P</td>
<td>Aireon Therapeutics, Inc.</td>
<td>CPAP Hydration Fluid. Pallet of Ultra-pure water specifically designed for CPAP usage. Pallet contains 96 cases, each case contains (24)-12 ounce bottles of CPAP Hydration Fluid. Using CPAC Hydration Fluid prevents growth of microorganisms (bacteria and fungus) which reduces respiratory health hazard, and increases equipment life which reduces CPAP lifecycle costs. Price includes UPS Ground shipping for customers in the Pacific Time Zone of the US</td>
<td>$5,428.28</td>
<td>1 Year</td>
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<td>A-20C</td>
<td>ABFA05</td>
<td>Axolotl Biologix, Inc</td>
<td>Amnion-Derived Ambient Allograft Liquid - .50ml, 1.5 Yr Shelf Life, Order Qty 1-20 units</td>
<td>$349.75</td>
<td>1 Year</td>
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<td>A-20C</td>
<td>ABFA10</td>
<td>Axolotl Biologix, Inc</td>
<td>Amnion-Derived Ambient Allograft Liquid - 1.0ml, 1.5 Yr Shelf Life, Order Qty 1-20 units</td>
<td>$567.84</td>
<td>1 Year</td>
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<td>SIN</td>
<td>Part #</td>
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<td>GSA</td>
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<tr>
<td>A-20C</td>
<td>ABFA20</td>
<td>Axolot Biologix, Inc</td>
<td>Amnion-Derived Ambient Allograft Liquid - 2.0ml, 1.5 Yr Shelf Life, Order Qty 1-20 units</td>
<td>$918.09</td>
<td>1 Year</td>
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<td>A-20C</td>
<td>ABM23</td>
<td>Axolot Biologix, Inc</td>
<td>Allograft Membrane Patch - 2cm x 3cm, 2 Year Shelf Life, Order Qty 1-20 units</td>
<td>$699.50</td>
<td>1 Year</td>
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<tr>
<td>A-20C</td>
<td>ABM44</td>
<td>Axolot Biologix, Inc</td>
<td>Allograft Membrane Patch - 4cm x 4cm, 2 Year Shelf Life, Order Qty 1-20 units</td>
<td>$1,224.12</td>
<td>1 Year</td>
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<td>A-20C</td>
<td>ADG02</td>
<td>Axolot Biologix, Inc</td>
<td>Axolot DualGraft™ 1cm x 2cm is a bi-layered dehydrated human amnion membrane allograft (dhAM) derived from the amniotic lining of the placenta. Axolot DualGraft™ can be used as a barrier and has properties known to advance soft tissue repair and reconstruction. Axolot DualGraft™ simplifies the application process by positioning the epithelial surfaces facing outwards, eliminating application placement limitations. Axolot DualGraft™ is classified as minimally manipulated under FDA regulation 21 CFR Part 1271 and section 361 of the PHS.</td>
<td>$641.94</td>
<td>1 Year</td>
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<td>A-20C</td>
<td>ADG06</td>
<td>Axolot Biologix, Inc</td>
<td>Axolot DualGraft™ 2cm x 3cm is a bi-layered dehydrated human amnion membrane allograft (dhAM) derived from the amniotic lining of the placenta. Axolot DualGraft™ can be used as a barrier and has properties known to advance soft tissue repair and reconstruction. Axolot DualGraft™ simplifies the application process by positioning the epithelial surfaces facing outwards, eliminating application placement limitations. Axolot DualGraft™ is classified as minimally manipulated under FDA regulation 21 CFR Part 1271 and section 361 of the PHS.</td>
<td>$1,095.08</td>
<td>1 Year</td>
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<td>Part #</td>
<td>Mfg</td>
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<td>A-20C</td>
<td>ADG16</td>
<td>Axolotl Biologix, Inc</td>
<td>Axolotl DualGraft™ 4cm x 4cm is a bi-layered dehydrated human amnion membrane allograft (dhAM) derived from the amniotic lining of the placenta. Axolotl DualGraft™ can be used as a barrier and has properties known to advance soft tissue repair and reconstruction. Axolotl DualGraft™ simplifies the application process by positioning the epithelial surfaces facing outwards, eliminating application placement limitations. Axolotl DualGraft™ is classified as minimally manipulated under FDA regulation 21 CFR Part 1271 and section 361 of the PHS.</td>
<td>$2,659.48</td>
<td>1 Year</td>
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<td>A-20C</td>
<td>ADG24</td>
<td>Axolotl Biologix, Inc</td>
<td>Axolotl DualGraft™ 4cm x 6cm is a bi-layered dehydrated human amnion membrane allograft (dhAM) derived from the amniotic lining of the placenta. Axolotl DualGraft™ can be used as a barrier and has properties known to advance soft tissue repair and reconstruction. Axolotl DualGraft™ simplifies the application process by positioning the epithelial surfaces facing outwards, eliminating application placement limitations. Axolotl DualGraft™ is classified as minimally manipulated under FDA regulation 21 CFR Part 1271 and section 361 of the PHS.</td>
<td>$3,182.74</td>
<td>1 Year</td>
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<td>A-20C</td>
<td>AS01</td>
<td>Axolotl Biologix, Inc</td>
<td>Axolotl Shot™, 1mL premeasured syringe of Axolotl Biologix an ambient temperature (25±2°C, 77±13°F) bioactive regenerative fluid, terminally irradiated for use in surgical applications. Axolotl Shot™ is derived from the amniotic components of the placenta to advance soft tissue repair and reconstruction. Axolotl Biologix uses our proprietary BioSym™ process to manufacture Axolotl Shot™.</td>
<td>$1,548.21</td>
<td>1 Year</td>
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<tr>
<td>A-20C</td>
<td>AS02</td>
<td>Axolotl Biologix, Inc</td>
<td>Axolotl Shot™, 2mL premeasured syringe of Axolotl Biologix an ambient temperature (25±2°C, 77±13°F) bioactive regenerative fluid, terminally irradiated for use in surgical applications. Axolotl Shot™ is derived from the amniotic components of the placenta to advance soft tissue repair and reconstruction. Axolotl Biologix uses our proprietary BioSym™ process to manufacture Axolotl Shot™.</td>
<td>$2,799.73</td>
<td>1 Year</td>
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</tbody>
</table>
Case containing (12) x 100-count boxes of individual DisposeRx packets is the perfect site-of-use solution for high-demand drug disposal environments such as pharmacies, hospitals, hospices, palliative care centers, nursing homes, assisted living facilities, and pain management clinics. Each DisposeRx packet when added to warm water in a pill vial containing unused medications renders the medication unusable and safe for household trash disposal. DisposeRx packets are non-toxic, biodegradable, environmentally friendly and are the only site-of-use drug disposal product that renders any medication -- capsules, liquids, pills or tablets -- non-retrievable preventing misuse, abuse and diversion – the three sources of new addictions, overdoses and deaths.

Kleiner Device Labs KG 1, Graft Delivery Complete System Kit. Each kit includes a patented graft delivery cannula, plunger, and attachable funnel reservoir for loading prepared bone graft material. Kits are sold and packaged in boxes of 6 individual sterile packs.

Kleiner’s KG 1 system provides a 21.0 cm length rectangular syringe barrel, plunger and attachable funnel for spinal surgical site access and graft material delivery. Kleiner’s patented design expedites minimally disruptive delivery of any flowable graft material to the intervertebral disc space. Its unique design allows 40% more efficient flow of bone graft material than the conventional, 8mm round-end dispensing cannula. It’s wedge tip allows it to enter collapsed disc spaces, while its two large lateral exit ports double the dispersion of graft, effectively eliminating jamming. The biportal extrusion design of the KG 1 ejects graft into the prepared area of the disc space bilaterally, instead of just at the anterior portion of the disk. This leaves a natural void for fusion cage placement.

Koelis, Inc. 3D Endocavity End-Fire Probe for Transrectal Application

Koelis, Inc. 3D Endocavity Side-Fire Probe for Transperineal Application

Koelis, Inc. Transperineal Application including TRUS Prostate Mapping (Promap software) and 3D Endocavity Side-Fire Probe
<table>
<thead>
<tr>
<th>SIN</th>
<th>Part #</th>
<th>Mfg</th>
<th>Description</th>
<th>GSA</th>
<th>Warranty</th>
</tr>
</thead>
<tbody>
<tr>
<td>A-94</td>
<td>KAPP-3000-R</td>
<td>Koelis, Inc.</td>
<td>Transrectal Application including TRUS Prostate Mapping (Promap software) and 3D Endocavity End-Fire Probe</td>
<td>$42,869.35</td>
<td>1 Year</td>
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<tr>
<td>A-94</td>
<td>KDNG00-96</td>
<td>Koelis, Inc.</td>
<td>Disposable Guides for 3D Endocavity End-Fire probe (96x units)</td>
<td>$3,055.28</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KIT-2049</td>
<td>Koelis, Inc.</td>
<td>Central Unit Computer Beam including software - V3</td>
<td>$29,788.94</td>
<td>1 Year</td>
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<tr>
<td>A-94</td>
<td>KIT-4011</td>
<td>Koelis, Inc.</td>
<td>Footswitch</td>
<td>$954.77</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KMNT-3000-SE</td>
<td>Koelis, Inc.</td>
<td>Trinity Annual Maintenance Program - SERENITY</td>
<td>$15,371.86</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KMRD-MR</td>
<td>Koelis, Inc.</td>
<td>MR Draw - MR Imaging Pack including MR Image Preperation and VM (Volume Measurements) software</td>
<td>$14,894.47</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KPHA.01.EL</td>
<td>Koelis, Inc.</td>
<td>STEADY PRO® complete set including probe holding arm with simple I and U extensions and STEADY PRO® support for 3D Endocavity Side-Fire Probe</td>
<td>$9,834.17</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KPHA.EC.2</td>
<td>Koelis, Inc.</td>
<td>STEADY PRO® support for 3D Endocavity End-Fire probe: cleanable external part (x5 units) and sterlisable internal parts of probe interface - Transperineal</td>
<td>$5,633.17</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KRNG.EL1.18-5</td>
<td>Koelis, Inc.</td>
<td>Reusable Mini Grid 18G Guides for Transperineal 3D Endocavity Side-Fire Probe (5x units)</td>
<td>$3,723.62</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KRNG.EL4.18-5</td>
<td>Koelis, Inc.</td>
<td>Reusable Full Grid 18G Guides for Transperineal 3D Endocavity Side-Fire Probe (5x units)</td>
<td>$6,206.03</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KSOF-2602-L</td>
<td>Koelis, Inc.</td>
<td>Promap 2nd Look software for Trinity</td>
<td>$19,286.43</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KSOF-2602-PET</td>
<td>Koelis, Inc.</td>
<td>Promap PET Software - PET &amp; CT Fusion for Trinity</td>
<td>$19,286.43</td>
<td>1 Year</td>
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<tr>
<td>A-94</td>
<td>KSOF-2602-PET-D</td>
<td>Koelis, Inc.</td>
<td>Promap PET Software - PET &amp; CT Images Preparation for MR Draw</td>
<td>$8,974.87</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KSOF-T-MR</td>
<td>Koelis, Inc.</td>
<td>MR Imaging Pack including MR Fusion and VM (volume measurements) software for Trinity</td>
<td>$22,055.28</td>
<td>1 Year</td>
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<tr>
<td>A-94</td>
<td>KTGP.18</td>
<td>Koelis, Inc.</td>
<td>Reusable Full Grid 18G Guides for Transperineal Application including Promap-GR software and hardware *for new Trinity units</td>
<td>$13,271.36</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-94</td>
<td>KURO-3000-2</td>
<td>Koelis, Inc.</td>
<td>TRINITY - 3D Prostate Suite and fully intregated system</td>
<td>$62,633.17</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-92</td>
<td>BATTBACK-1</td>
<td>TruMed Systems, Inc.</td>
<td>Additional Battery back-up pack (extends battery backup from 9 hours to 18 hour guaranteed) Shipping to CONUS</td>
<td>$1,557.79</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-92</td>
<td>GLDYr-1</td>
<td>TruMed Systems, Inc.</td>
<td>Annual Gold Service Package , Silver Service Package Plus, Customer Service with 24/7 access Onsite service with 4 hour response (location dependent), Vaccine Insurance</td>
<td>$4,944.72</td>
<td>1 Year</td>
</tr>
<tr>
<td>SIN</td>
<td>Part #</td>
<td>Mfg</td>
<td>Description</td>
<td>GSA</td>
<td>Warranty</td>
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<tr>
<td>A-92</td>
<td>PatientMode</td>
<td>TruMed Systems, Inc.</td>
<td>Patient Mode Software Upgrade annual license. Provides the opportunity to track the dose by patient identifier</td>
<td>$988.94</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-92</td>
<td>PLATy-1</td>
<td>TruMed Systems, Inc.</td>
<td>Annual Platinum Service Package , Gold Service Package Plus, Onsite service with 2 hour response (location dependent), Vaccine Insurance</td>
<td>$6,211.06</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-92</td>
<td>PracticeInv</td>
<td>TruMed Systems, Inc.</td>
<td>Practice wide inventory (per site) annual. Allows single overview of inventory in non-AccuVax storage devices at the location</td>
<td>$1,242.21</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-92</td>
<td>ReOrder</td>
<td>TruMed Systems, Inc.</td>
<td>Reordering service (automatically reorder) based on par levels monthly (will require integration work prior to availability for DoD)</td>
<td>$627.14</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-92</td>
<td>SLVYr-1</td>
<td>TruMed Systems, Inc.</td>
<td>Annual Silver Service Package , Repair or Replace Warranty (up to 60 months from install), Customer Service with 8 AM - 5 PM access, Onsite service with 6 hour response (location dependent), AccuVax Software license , Full Alerting Suite, Portal Subscription, Probe calibration (as needed)</td>
<td>$3,738.69</td>
<td>1 Year</td>
</tr>
<tr>
<td>A-92</td>
<td>T1000-Purch</td>
<td>TruMed Systems, Inc.</td>
<td>AccuVax Hardware Kiosk (requires a service package) Robotic vaccine storage and dispensing device, Dimensions of 30.0&quot;W x 28.5&quot;D x 65.0&quot;H, 600 refrigerated storage locations, Refrigerated Drawer for up to 300 additional doses or other meds, 119 frozen locations, 7 Integrated NIST Calibrated Digital Data Loggers, Ethernet/wireless/LTE Connectivity, Biometric and/or password protected login, Battery Backup (up to 15 hours, 9+ guaranteed), Installation and Shipping to CONUS, Staff initial training and setup</td>
<td>$24,723.62</td>
<td>1 Year</td>
</tr>
</tbody>
</table>