

TRADEMARK ASSIGNMENT

Electronic Version v1.1
 Stylesheet Version v1.1

SUBMISSION TYPE:		NEW ASSIGNMENT	
NATURE OF CONVEYANCE:		RELEASE BY SECURED PARTY	
CONVEYING PARTY DATA			
Name	Formerly	Execution Date	Entity Type
J.P. MORGAN EUROPE LIMITED		05/27/2011	private limited company: UNITED KINGDOM
RECEIVING PARTY DATA			
Name:	SYSTAGENIX WOUND MANAGEMENT (US), INC.		
Street Address:	400 Crown Colony Dr., Suite 302		
City:	Quincy		
State/Country:	MASSACHUSETTS		
Postal Code:	02169		
Entity Type:	CORPORATION: DELAWARE		
PROPERTY NUMBERS Total: 1			
Property Type	Number	Word Mark	
Registration Number:	1928621	REG GRANEX	
CORRESPONDENCE DATA			
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Signature:	/Erin Frazier/		

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REEL: 004594 FRAME: 0409

Date:

07/28/2011

Total Attachments: 26

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**PARTIAL RELEASE OF SECURITY INTEREST UNDER SENIOR FACILITIES
AGREEMENT
AND TRADEMARK AND PATENT SECURITY AGREEMENTS**

This PARTIAL RELEASE OF SECURITY INTEREST UNDER the SENIOR FACILITIES AGREEMENT, the TRADEMARK SECURITY AGREEMENT and the PATENT SECURITY AGREEMENT (this Partial Release) is executed as of May __, 2011 by J.P. MORGAN EUROPE LIMITED in its capacity as Security Agent (the Security Agent).

WHEREAS, SYSTAGENIX WOUND MANAGEMENT (US) INC. (Systagenix US) and the Security Agent are (among others) parties to that certain Senior Facilities Agreement dated as of 26 NOVEMBER 2008, as amended as of 22 DECEMBER 2008, 14 July 2010 and 31 December 2010 (as the same may be further amended, supplemented or otherwise modified, renewed or replaced from time to time), (the Senior Facilities Agreement);

WHEREAS, pursuant to the Senior Facilities Agreement, Systagenix US executed and delivered to the Security Agent that certain Trademark Security Agreement dated as of January 13, 2009 and recorded with the United States Patent and Trademark Office on January 30, 2009 at Reel/Frame 3927/0148 ("Trademark Security Agreement");

WHEREAS, pursuant to the Senior Facilities Agreement, Systagenix US executed and delivered to the Security Agent that certain Patent Security Agreement dated as of January 13, 2009 and recorded with the United States Patent and Trademark Office on January 30, 2009 at Reel/Frame 022177/0037 ("Patent Security Agreement");

WHEREAS, pursuant to the Senior Facilities Agreement, the Trademark Security Agreement and the Patent Security Agreement, Systagenix US has granted to the Security Agent a continuing security interest in, *inter alia*, the Released Property; and

WHEREAS, Systagenix US has requested that the Security Agent release the liens and security interests granted to the Security Agent by Systagenix US (pursuant to the Senior Facilities Agreement, the Trademark Security Agreement and the Patent Security Agreement) in Systagenix US's right, title and interest in and to the Released Property defined below (to the extent of Systagenix US's interests therein).

NOW, THEREFORE, for good and valuable consideration, receipt of which is hereby acknowledged:

1. Definitions. Capitalized terms used herein and not otherwise defined herein shall have the meanings given such terms in the Senior Facilities Agreement. For purposes of this Partial Release, "Released Property" shall mean all right, title and interest in and to the Transferred Assets (as defined below in Exhibit A) described in Exhibit A attached hereto.
2. Partial Release. The Security Agent hereby:
 - a) releases the liens and security interests granted by Systagenix US to the Security Agent pursuant to any security agreement (including but not limited to the Senior Facilities Agreement, the Trademark Security Agreement and the Patent Security Agreement) in the right, title and interest of Systagenix US in and to the Released Property as defined herein to the extent of Systagenix US's interest therein; and

b) to the extent the Security Agent shall be deemed to have any right, title or interest in the Released Property, retransfers and reassigns to Systagenix US all of such right, title and interest solely with respect to the Released Property as defined herein.

3. Full Force and Effect. Except as expressly modified hereby, the Senior Facilities Agreement, the Trademark Security Agreement and the Patent Security Agreement shall remain in full force and effect in accordance with the provisions thereof on the date hereof.

4. Governing Law. THIS PARTIAL RELEASE SHALL BE GOVERNED BY THE LAWS OF THE STATE OF NEW YORK AND THE APPLICABLE FEDERAL LAWS OF THE UNITED STATES OF AMERICA, WITHOUT REGARD TO ANY CONFLICT OF LAWS PRINCIPLES THAT WOULD RESULT IN THE APPLICATION OF THE LAWS OF ANOTHER JURISDICTION.

[Signature page to follow]

IN WITNESS WHEREOF, this Partial Release has been duly executed as of the date first written above.

J.P. MORGAN EUROPE LIMITED
as Security Agent

By: 

Name: _____

Title: Jonathan Richards
Executive Director

{Signature Page to Partial Release of Security Interest}

DN1/60186-922

Exhibit A

Transferred Assets

Definitions

As used herein, the term "Product" means a certain prescription gel sold under the trade name "Regranex[®]."

As used herein, the term "Business" means the business of manufacturing, selling and distributing the Product, and expressly excludes any other business or product line of Systagenix, or any of its Affiliates.

As used herein, the term "Transferred Assets" means those assets, properties and rights relating to the Product and Business, as described below.

As used herein, the term "Systagenix" means Systagenix Wound Management, LTD.

Transferred Assets

- (i) Inventory. All Inventory, including without limitation, inventory of finished Product, Product specific work in process and all becaplermin ("API") and raw materials, packaging and supplies used primarily in connection with the manufacture of the Product, wherever located, in transit or otherwise;
- (ii) Equipment. The machinery, equipment, tools, hardware, spare parts, furniture, furnishings, fixtures, office supplies, computers and peripherals and other tangible personal property identified on Schedule 1 (collectively, "Transferred Equipment");
- (iii) Business Records. The following records and files to extent primarily relating to the Business, the Product or other Transferred Assets and in possession of, or reasonably available to, the Systagenix or a Systagenix Affiliate (and all copies and tangible embodiments of the foregoing (in whatever form or medium)); and, to the extent such records or files are not reasonably separable from documents or databases that do not relate to the Business, the Product or other Transferred Assets, copies of such records or files (redacted as to any information not relating to the Business, the Product or other Transferred Assets): (1) vendor lists, (2) customer lists, (3) pricing lists for the Product, (4) advertising, marketing, sales and promotional materials, (5) patterns, plans, designs, research data, formulae and manufacturing processes, quality testing procedures, quality control records, operating manuals, drawings, manuals, data, records, procedures, research and development records, design history files, compositions, drawings, specifications, improvements, proposals, technical and computer data and programs, and related documentation and (6) such other material records primarily relating to the Business, the Product or other Transferred Assets, to the extent such other records are permitted to be transferred under applicable Law;
- (iv) Intellectual Property. (A) The Trademarks and the Patents set forth on Schedule 2 and all IP Rights relating thereto, (B) all IP Rights under the Transferred Contracts and (C) all other IP Rights (other than Internet domain names, which are described in paragraph (v) below) used, or held for use, exclusively in connection with the Product or the Business as currently conducted (collectively, the "Transferred IP");

(v) Telephone Lines, Websites and Domain Names. Telephone numbers (including all rights in customer telephone service lines), websites (including website content and copyrightable work displayed thereat) and domain names primarily related to the Business, which are set forth on Schedule 3 and all IP Rights relating to such websites and domain names;

(vi) Contracts. Each of the Transferred Contracts listed on Schedule 4 and all rights thereunder from and after the Closing;

(vii) Goodwill. The goodwill generated by or associated with the Business and the Product;

(viii) Claims. All claims, causes of action and rights of recovery, known or unknown, absolute or contingent, relating to any Transferred Asset (including rights under any warranties, representations and guarantees made by suppliers, manufacturers, contractors or others in connection with the operation of the Business or affecting the Transferred Equipment or the Inventory) and all counterclaims, set-offs or defenses that may exist with respect to any Assumed Liability; and

(ix) Derivative Rights in Transferred IP. (A) All income, royalties and other payments due or payable with respect to the Transferred IP, (B) the right to sue for and obtain remedies against (including the collection of damages) past, present and future infringement, misappropriation or dilution with respect to the Transferred IP, (C) rights of priority and protection of interests under applicable Laws with respect to the Transferred IP and (D) rights to defend against claims that any of the Transferred IP infringes the intellectual property rights of any third party.

Schedule 1
Transferred Equipment

See attached.

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Asset List

Work Instruction	
Status: Effective	Effective Date: 8/5/2018 16:42:03 (EST)
Title: Site Validation Program	
Doc. No. P884-DOC-8892	Version: 18.0, CURRENT

Attachment 5: Computerized Systems Inventory

System Name	Location	Identifier	Description
Honeywell Smart Building Management EBI System Program Area	Regrex Area, Building 1	4	The new Honeywell Enterprise Building Integrator (EBI) application version 310.1 is a graphic configured application developed to interface with the existing Direct Digital Control (DDC) Panels, which controls and monitors the HVAC System of the Regrex Manufacturing Area. The EBI application version 310.1 resides on a Dell Server PowerEdge SC7400/Intel Xeon processor, running on a Windows 2008 OS Server Standard Edition with a Microsoft SQL Server 2008 database application, interfacing with the RT844 Excel Plus Controllers (DDC Panels) through a Building Network Adapter (BNA) bus type interface. The Honeywell EBI version 310.1 allows the operator to establish and monitor parameters for temperature, relative humidity, differential pressure and alarm limits. The control system executes functions such as command valve operations, monitoring established parameters in rooms, alarms and actual airflow (CFM). The DDC panels receive the rooms condition data (Temperature, Humidity and Pressure) from the sensors and transmitters, and with the support of the EBI version 310.1, the data is processed in order for the DDC to actuate our valves and dampers to maintain the desired parameters previously set by the operator.
Honeywell Controller Stability/Reserve Samples Area	Room 106 Building 1	2a	Stand alone closed controller used to maintain the temperature and relative humidity conditions of the Stability/Reserve Samples Area. An independent chart recorder inside the Stability/Reserve Samples room records the temperature and relative humidity conditions on a seven day range.
CalPro Software Application version 3.0.1b	Metrology Area, Building 1	3	Computer-Related Calibration System (CAL.pro) is a stand alone closed system composed of the following: IBM NetVista Personal Computer, Iomega Zip Drive backup device, Hewlett Packard LaserJet printer, Libart UPS/Inverter tower, SQLbase database platform, Windows 2000 operating system platform and McAfee VirusScan, version 4.6.1 S/P1. This system keeps an inventory of regular and reference instruments, automatically determines when they are due for a calibration check, generates their calibration work orders and documents the calibration results. The system is capable of generating reports based on transaction queries to the SQLbase database.

Work Instruction	
Status: Effective	Effective Date: 8/5/2010 16:42:03 (EST)
Title: SRA Validation Program	
Doc. No.: PEGA-CCC-5622	Version: 18.0, CURRENT

Attachment S: Computerized Systems Inventory (continuation)

System	Area of Service	Classification as per PEGA, DEC-1978	System Description
Mesimo Software Application version 4.1.1	Metrology Area, Building 7	3	Mesimo is a configurable, client/server or terminal/server software package for the automated management of maintenance activities. It is used to acquire, analyze, report and store data generated. The hardware architecture for the system is a Compaq ProLiant Server in which both the application and the database reside. The clients are IBM PCs connected to the server through the existing LAN infrastructure using a V-LAN configuration.
Waters Millennium ³² Data Acquisition System version 4.0	CO Analytical Laboratory	4	The Waters Millennium ³² software version 4.0, is an off the shelf configurable software application that is designed to control, acquire data and report results from chromatographic instrumentation. It incorporates an Oracle database to organize information. The application is installed on a network using Ethernet protocol, including a server, client PCs and LADIE ³² s. The database server contains the database and provides central security and information storage for the system. The client PC provides the user access to the system and also performs data analysis, but no data is stored there. The LADIE ³² , also called an acquisition server, provides the connection between the chromatographic instruments and the Millennium ³² system.
Elektronik Controllers Compressed Air System	Engineering/Utilities	2a	The Elektronik micro-processor non-configurable firmware regulator serves as the controller of the air buffer/loaded sequencing functions of each such as running parameters, alarms and safety indicators. The system operates as a standalone closed system, although the controller has the capability of external communication through the use of field bus systems and/or Ethernet communication.
Controllers Air Dryers and PLC-1502	Engineering/Utilities	2a	The air dryer timer board is considered a controller with non-configurable firmware, which operates as a closed system. As part of the dual-air dryer configuration, the air dryer timer board controls the time period on which one tower is drying the main air stream, while the second tower is being regenerated by a purged air stream. The timer boards are independent units with no interface between them. Each timer board is considered as a stand-alone system. The timer board contains a standard control panel with voltage selection and DIP switches, to select among standard or high-pressure units, operating cycles, test mode and purge accumulator settings. Also, it contains operational warnings and alarm light indicators.
Regranex Manufacturing Area Process Control System	Regranex Manufacturing	5	The Regranex Process Control System controls the manufacturing processes and monitors the utilities in the Regranex Manufacturing Area. The Main PLC S42E is linked via Ethernet communication to the Allen Bradley SLC's processors as part of the WFI SWS, Pure Steam Generator, Buffer Preparation TCM and Gel Preparation TCM. The CIP PLC S20E also is connected to the Main PLC through Ethernet communication. The existing consists of three (3) workstations in a LAN configuration connected through a switch to the Main PLC, which is connected to all the field devices. The workstations are located in the Buffer Preparation Room, Gel Preparation Room and Utilities Room. A HP Compaq server is installed to support security and data backup features.

Work Instruction	
Status: Effective	Effective Date: 5/5/2010 10:42:03 (EST)
Title: Sba Validation Program	
Doc. No. PEGA-DCG-0002	Version: 15.0, CURRENT

Attachment 5: Computerized Systems Inventory (continuation)

System Name	Location	Count	Description
Regrenex USP Water System Control System	Engineering Utilities	8	The SCADA/FLO Network Infrastructure consists of the following main components: SCADA/FLO System (also identified as CP-5): An integrated system implemented for the monitoring activities of the USP Water Separation System (US Filter PreVIEW 100), the automatic controlling of the pre-treatment pumps and the USP Water Distillation System, including the Finished Water Storage Tank, distillation pumps, heater/cooler heat exchangers, supply instruments (TDS analyzer, Conductivity meter, pH meter, mass flow meter, temperature and pressure indicators) and return instruments (TDS analyzer, Conductivity meter, pH meter, mass flow meter, temperature and pressure indicators). CP-5 will also have the capability of monitoring the existing Regrenex USP Water Distribution Loop through an interface with the existing Regrenex SCADA Production Management System.
Keys Validator 2000 Units, Software version 2.21 with PC Laptops	QA Validation Department	3	This is a standard thermal validation system, which can be operated in standalone mode or through the use of the Validator 2000 software package installed on a laptop PC. The Validator 2000 Software version 2.21 is a standard application software, which is used with the Validator 2000 Data Logger hardware system for the monitoring and documentation of critical parameters in thermal processes. The system provides functionalities for calibrating sensors, performing qualification studies, calibration verification and generating reports. The system is capable to store electronic data from the calibration and qualification runs executed. Current practices at CMJ considered the hardcopy of the record as the official document in which the company relies on.
Control System of the Pure Steam Generator (PK-791) Model 500-S-1	Regrenex Utilities Area	4	The Pure Steam Generator Model 500-S-1 is designed to produce pyrogen-free pure steam from feed water of USP quality. Its structure consists of one column, pre-heater and other peripheral components. The control system of the PEG consists of an Allen Bradley PLC 505 (Small Logic Controller), Allen Bradley I/O Modules, a Dr-485 Link Coupler and a control panel which consists of an alarm indicator panel, power ON/OFF button, alarm reset button, Newport 02 pressure receiver, a Polymatron Model 5823 Quality Analyzer, and a Yokogawa UR1000 recorder (for reference only). The control system, transformer(s), fuses, modules etc., are mounted within a control box, which is integral to the unit's frame. This system controls and monitors the steam generator operation based on predefined parameters programmed into the PLC controller. The Pure Steam 500-S-1 Pure Steam Generator is used to generate steam for cleaning and sterilization processes in the Regrenex Area at Building 1.

Work Instruction	
Status: Effective	Effective Date: 6/6/2010 15:42:03 (EST)
Titer Site Validation Program	
Doc. No. PSGA-DOC-0022	Version: 15.0, CURRENT

Attachment 8: Computerized Systems Inventory (continuation)

System	Area or Facility	Classification (FDA 21 CFR 312.62)	System Description
Control System of the Finn Aqua Water Still Generator Model 200-S-6	Regenerex Utilities Area	4	The Finn-Aqua multiple-effect Water Still Model 200-S-6 is designed to produce pure, pyrogen-free distilled water from preheated feedwater of USP quality by evaporating and condensing the feedwater, at the Regenerex Area. Its structure consists of five (5) columns, pre-heaters, condenser and other peripheral components. The control system consists of an Allen Bradley PLC 5/30 (Smart Logic Controller) with Ethernet coupler to communicate with the Regenerex Main PLC 5MDE, Allen Bradley I/O Modules, a DH-485 Link Coupler and a control panel consisting of an alarm indicator panel, temperature monitoring panel, sequence stop panel, emergency stop panel, feed water pump control switch, main power switch, Contraves Message Center display for system parameters, Polymatron Morse 8020 Quality Analyzer, Pomegran PC-1520 Temperature Analyzer and a Yokogawa CR1000 recorder. This system controls and monitors the water still operation based on predefined parameters programmed into the PLC controller.
Controller for the Finn Aqua Autoclave 104 Model 121512-D-5-511	Regenerex Manufacturing	2b	The control system components are the following: an Autonics II Microterminal AL1001, power supply module, CPU, Analog and Digital I/O module, Digital I/O module, a dot matrix printer and a chart recorder. This standalone and cabled system has the capability of connecting to an external device in terms of reporting temperature and pressure parameters, but this capacity is restricted due to the configuration of the hardware. This type of system is considered as a single-point system in which a routine environment and process parameters are used to comply with a determined manufacturing process.
Nordson 2002 Tube Filler Machine	Regenerex Packaging Building 1	2b	The Nordson 2002 Tube Filler Machine is a high-speed packaging machine. Tubes are automatically pushed along with pellets into their carton. Locks can either be dispensed from another magazine or from a rail. Finally the machine will automatically tack the carton shut and emboss the end of the cartons with the lot number and expiration date. The control system comprises safety functions as well as check functions. The control panel is provided with push buttons, indicator lamps and switches in order to allow the machine operator to control the different machine function settings. This is a controller with non-configurable hardware.

Work Instruction	
Status: Effective	Effective Date: 2/5/2010 10:42:03 (EST)
Title: Site Validation Program	
Doc. No. PSGA-000-0005	Version: 10.0, CURRENT

Attachment 5: Computerized Systems Inventory (continuation)

System Name	Department	Asset ID	Description
Nordenmatic 2002M Tube Filler and Capper	Regenerex Manufacturing	2a	The Nordenmatic 2002 is a high-speed tube-filling machine for an output up to 200 tubes per minute. The control system completes safety functions as well as check functions. The Norden Control System is easily programmable from the operating panel of the machine. The control panel is provided with push buttons, indicator lamps and switches to allow the machine operator to control the different machine function settings. This is a controller with non-volatile memory.
Vitek Identification System	Micro Lab	3	The Vitek system version R30-01 is an automated system designed to identify gram negative/gram-positive bacteria and yeast. The Vitek System consists of test cards, a flowcell, a reader/analyser, a PC workstation with the industrial bioMARDON SM Software with Vitek VTK software version R30-01, and a Printer. The computer processes results, collecting them from the reader/analyser, storing them during the incubation period, transmitting interim results to the screen or printer, and automatically sending final results to the printer. The system is being used as a stand-alone system generating printed reports that are recorded on laboratory notebooks.
Micro Lab Spreadsheets	Micro Lab	5	The spreadsheets use using Microsoft Excel 2000, and MikLab Station Software. They reside in the San German data share drive and can be accessed with an authorized user account from any computer on the network. The spreadsheet files are used to keep records of the results for the sampling testing process for the following areas: Environmental, Sampling, Personnel Qualification, Product Testing, Media Fill, Special Testing, Utilities. The spreadsheets have columns with formulas to keep track of the number of observations that fall inside/outside of acceptance levels, columns for the date, classification of the organism, indication if a investigative report needs to be generated or not, and indication of the sample method. They also contain graphical representations of the results.

Work Instruction	
Status: Effective	Effective Date: 5/5/2010 10:42:03 (EST)
Title: EMS Verification Program	
Doc. No.: PSGA-DOC-0052	Version: 1.0, CURRENT

Attachment 5: Computerized Systems Inventory (continuation)

System Name	Area	Classification	System Description
Stability Program	Stability and Reserve area	4	The Stability Program Spreadsheet is used to keep records of the stability lots and their testing necessary time intervals. Inside the spreadsheet, there are several tabs to record the different products stability lots. As a spreadsheet, and based on the lot manufacturing data, it calculates the time intervals (3, 5, 8, 12 and 15 months) for necessary lot testing. Also, a formula or function defined as "trigger" allows the user to automate the classification of the results. The Stability Program Spreadsheet is accessed through the Puerto Rico Pharma MSN-CDM-San German directory on the file server (L) server. The spreadsheet runs using Microsoft Excel 2003, on a Windows 2000 Platform. The access to the folder is controlled by the NCB network user name and password. The network accounts are assigned permissions to access the folders. The access to the folder will allow to 'read' or 'write' to the spreadsheet depending on the privileges of the account. Without access, the person receives an access denied message when trying to access the folder.

Work Instruction	
Status: Effective	Effective Date: 5/5/2010 16:42:05 (EST)
Title: SAs Validation Program	
Doc. No. FSGA-DCG-0852	Version: 16.0, CURRENT

Attachment B: Qualified Equipment List
Note: Computerized Systems are listed as part of Attachment #5

Doc. No.	Equipment	Area of Service	Description
0878	T Transfer Line	Room 232, Filling Area, Bldg. 1	Stainless steel 316 electropolished piping line used to transfer the gel product from the Portable Tanks located in Room 234 (In-Process Cold Room) to Filling Room 232.
0780	Manual Cleaning (Compounding / Filling Equipment)	Clean at Room 243 Washing Area, Bldg. 1	Equipment components manually cleaned, which are part of the Regranex Manufacturing Equipment.
0365	Plus Aqua Autoclave 134 Model 121512B (S/N: 88596)	Room 243 Washing Area, Bldg. 1	Clean steam autoclave used for the sterilization of components and equipment used during the manufacturing process of Regranex.
0598, 0367	Buffer Vessel (including expansion Tank T-206)	Room 255, Buffer Preparation, Bldg. 1	Equipment used to prepare the buffer solution used as part of the Regranex Manufacturing Process.
0337, 0338, 0339, 0340, 0341	Flash Blend and Distribution System (pre-filters, pump and hoses)	Room 255, Gel Formulation, Bldg. 1	The Flash Blend and Distribution System is used during the NaOH transfer to a stream of buffer solution, which is pumped, sterile filtered and transfer into the Gel Free Turboemulsifier.
0548	Portable Tanks (A to H)	Room 255 Gel Formulation; Room 234 In-process Cold Room, Bldg. 1	Stainless steel portable tanks used to hold the gel product after completing the gel manufacturing process, during the storage inside the In-process Cold Room and used during the filling process.
0343, 0344, 0345, 0347	Disinfectant Tank T-301, Expansion Tank T-302, Vent Filter and Pump	Room 255 Gel Formulation, Bldg. 1	Homogenizer equipment used during the manufacturing process of Regranex Gel 0.01%.
0316, 0313	Active Drug Assembly (Addition Pot T-104, Pre-filters T-105-A/B, Filling Vessel T-101, WPI Chitos Pol S-103)	Room 255 Gel Formulation, Bldg. 1	Equipment used to hold and transfer the active drug (Benzopermin) to the Ross Turboemulsifier.
0742	Puffer Hubbard Freezer (Model UP3050A-16, S/N: 010M-082228-1M)	GMJ Freezers Room, Warehouse Area, Bldg. 7	Freezer with an operational range down to -80 °C, which is used to freeze the Regranex Gel Packs.
0743	Rever Upright Freezer (Model UL7050A-14, S/N: P225-20178U-PE)	GMJ Freezers Room, Warehouse Area, Bldg. 7	Freezer with an operational range down to -80 °C, which is used to freeze the Regranex Gel Packs.
0340	Puffer Hubbard Upright Freezer (Model SLT-23V-40A34, S/N: V28K48085W1C)	GMJ Freezers Room, Warehouse Area, Bldg. 7	Freezer with an operational range of -10 to -40 °C, which is used to freeze the Regranex Gel Packs.
0341	Puffer Hubbard Upright Freezer (Model SLT-23V-40A34, S/N: V28K48085W1C)	GMJ Freezers Room, Warehouse Area, Bldg. 7	Freezer with an operational range of -10 to -40 °C, which is used to freeze the Regranex Gel Packs.
0332	Puffer Hubbard Freezer (S/N: R17C-534-035-RC)	Warehouse Area, Bldg. 7	Freezer with an operational range of -80 to -85 °C.
0300	Wet-In RF-07 Refrigerator	Room 172, Building 1	Refrigerator with an operating range of 2-8 °C.

Work Instruction	
Status: Effective	Effective Date: 5/5/2010 16:42:03 (EST)
Title: S&S Validation Program	
Doc. No. PESA-DCC-0082	Version: 15.0, CURRENT

Attachment 5: Qualified Equipment List (continuation)

Equipment ID	Equipment Name	Area of Control	Description
0708	Walk In Refrigerator RF-16	Warehouse Area, Bldg 7	Walk-in refrigerator with an operating range of 2-8 °C.
0811	Thaw Refrigerator RF-11 (S/N: P14E201008-PE)	Room 240, Thawing Area, Bldg. 1	Refrigerators used during the thawing process of Bacoplermin, operating range 2-8 °C.
0812	Thaw Refrigerator RF-12 (S/N: P14E201004-PE)	Room 240, Thawing Area, Bldg. 1	Refrigerators used during the thawing process of Bacoplermin, operating range 2-8 °C.
0890	Puffin Hubbard Freezer (Model RFB25-A14; S/N: Y12G-00725-VG)	Active Drug Storage Room 239, Building 1	Freezer with an operational range of -80 to -90 °C, used for the storage of active ingredient within the Regeneron manufacturing area.
0892	Puffin Hubbard Freezer (Model UF7018-A12; S/N: P18E-201114-PE)	Active Drug Storage Room 239, Building 1	Freezer with an operational range of -80 to -90 °C, used for the storage of active ingredient within the Regeneron manufacturing area.
0808	In Process Cold Room RF-16	Room 239, Building 1	Refrigerator-room with an operational range of 2-8 °C used for the in-process storage of Regeneron Gel water to the filling operations.
0890	Puffin Hubbard Freezer (Model RFB25-A12; S/N: P18E-201113-PE)	Warehouse Area, Bldg 7	Freezer with an operational range of -80 to -90 °C.
0847	Haeco Elite UltraLow Temperature Freezer (Model ULT1240-6-A12; S/N: UDF-000228-UF)	Room 198, Bldg 1	Freezer with an operational range down low to of -80 °C.
0808	RF-14 Refrigerator	Warehouse Area, Bldg 7	Walk-in refrigerator with an operational range of 2-8 °C.
N/A	Water System	Regeneron Utilities Area, Bldg 1	The OMI water purification system consists of a series of process treatments to produce four different qualities of water: Potable Water, Soft Water, USP Purified Water and Water for Injection. After passing through the pre-treatment equipment (chemical injection systems, Multimedia, Softener, Carbon filter, pre-filter, UV light, UV, RO unit, ODI unit, second UV and final filter), the water is sent to a Purified Water Storage Tank and pump through the distribution loop and to the different points of use. Water for Injection is obtained by distilling purified water using Finn Aqua multi-effect stills. There is a loop of WFI at the site, for the biological (Regeneron) production area. WFI is used for the following: Process water in biological (Regeneron) products; and equipment cleaning in the compound formulation area.
N/A	Steam System: Finn Aqua Pure Steam Generator 600-S-1	Regeneron Utilities Area, Bldg 1	Generator used to produce clean steam to support the operation of the Sterilization in Place (SIP) circuits and autoclave operation.

Work Instruction	
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Title: Site Validation Program	
Doc. No. FEGA-DOC-0002	Version: 16.0, CURRENT

Attachment 8: Qualified Equipment List (continuation)

Tag No.	Equipment	Area of Service	Description
N/A	HVAC System	Ceiling Area, Building 1	The Regranex HVAC System consists of 3 Air Handling Units (AHU), a reheat system, and a chiller system equipped with pumps to maintain the environmental conditions of the manufacturing rooms in terms of temperature, relative humidity and differential pressure, as applicable.
N/A	Dust collector DC-102 (JTD System)	Regranex Manufacturing Area, Bldg 1	Dust collection system (PulsePak Design 6, continuous Guly pulse jet cartridge dust and fume collector) supporting the Buffer and Weighing Areas.
N/A	Compressed Air System	Utilities Area, Building 1	The compressed air system consists of 2 Atlas Copco Oil-Free Air compressors, adsorption dryers, surge tanks as well as piping and valves. Compressed Air is used as part of the drying cycles of the CR runs as well as to support the pneumatics instrumentation used at the area.
0309	Nordenmatic Filler/Sealer	Room 202 Filling Room	The Nordenmatic 2002 is a high-speed tube-filling machine for an output up to 200 tubes per minute. It is provided with push buttons, indicator lamps and switches to allow the machine operator to control the different machine function settings.
0310	Nordenmatic Tube Feeder	Room 160, Tube Feed Room	The Norden Automatic Tube Feeder is a high speed machine that automatically picks and removes tubes from opened boxes, places the tubes onto a rigid tube transfer belt, and feeds them into the filling and capping machines.
N/A	Nordenmatic Cartoner	Room 224, Packaging Area	The Nordenpac 2002 Tube Cartoner Machine is a high-speed packaging machine. Tubes are automatically pushed along with feeders into their carton. Leaflets can either be dispensed from another magazine or from a roll. Finally the machines will automatically heat the carton spout and emboss the end of the cartons with the lot number and expiration date.
0307	Laminar Flow Hood (LFH)	Room 240, Thawing Area, Bldg. 1	Equipment used during the thawing/packaging process of Rosuvastatin.
0301, 0313, 0315, 0320, 0321, 0322, 0325, 0326, 0328, 0342	CIP System (Heat Exchangers, Detergent Neutralization Tank, PK-501, Return/Supply/Drain/ Surge/ Cassette Injection/ Acid Injection Pumps)	Room 244, Pre-wash Room	The CIP Cold System is used to clean the following manufacturing equipment and associated piping: Buffer Tank, Active Drug Assembly, Race Turbo-Emulsifier, Flashblend Line, Hot Transfer Line and Portable Tanks. The CIP Cold automatically sequences and delivers a series of wash and rinse solutions to the system to be cleaned. Flow and step duration rates are adjusted specifically for the equipment circuit being cleaned.

Work Instruction	
Status: Effective	Effective Date: 5/5/2010 16:42:03 (EST)
Title: Site Validation Program	
Doc. No. PEGA-DOC-8882	Version: 1.0, CURRENT

Attachment B: Qualified Equipment List (continuation)

Tag No.	Equipment	Area of Service/Location	Description
0235	Precision Scientific Incubator R-22 (Model 30 MP, S/N: 867030102)	Microbiology Laboratory Area, Room 183 Bldg 1	Incubator with an operational range of 30 to 35 °C.
0725	Precision Scientific Incubator (Model 31030, S/N: 896011066)	Microbiology Laboratory Area, Room 183 Bldg 1	Incubator with an operational range of 20 to 25 °C.
0395	Precision Scientific Incubator (Model 30M, S/N: 897030105)	Microbiology Laboratory Area, Room 181 Bldg 1	Incubator with an operational range of 30 to 35 °C.
0394	Fisher Scientific Isotemp Plus Laboratory Refrigerator (Model 452250A12, S/N: 2305-222036-02)	Microbiology Laboratory Area, Room 181 Bldg 1	Refrigerator with an operational range of 2 to 6 °C.
0682	Precision Scientific Incubator (Model 30 MP, S/N: 896030602)	Microbiology Laboratory Area, Room 183 Bldg 1	Incubator with an operational range of 20 to 25 °C.
0275	Fisher Chest Freezer (Model BC1010, S/N: 880702055)	Microbiology Laboratory Area, Room 183 Bldg 1	Freezer with an operational range of -20 +/- 4 °C.
0277	Heraeus Incubator (Model B30, S/N: 30000025)	Microbiology Laboratory Area, Room 216 Bldg 1	Incubator with an operational range of 55-60 °C.
0390	Fisher Isotemp Plus Laboratory Refrigerator (Model 4525-A12, S/N: P125-333337)	Microbiology Laboratory Area Room 182, Building 1	Refrigerator with an operational range of 2 to 6 °C.
0599	Vitek	Microbiology Laboratory Area, Room 219 Bldg 1	The Vitek Microorganism Identification System consists of test cards, a Filter/Sealer, a Reader/Incubator, a PowerPC workstation with the bioMERSION™ Vitek VTK software, and a printer. A colorimeter single-beam photometer is used during the proper inoculation of the test cards. The Filter/Sealer uses vacuum to fill the test card and seal it. The Reader/Incubator holds the prepared test cards in a temperature controlled environment, and scans them with photometric sensors, searching for changes in color and turbidity, which reports to the computer. Comparing the microorganism's biochemical reactivity pattern to profiles in the system's database makes identification.
N/A	Fisher Scientific pH Meters (1 unit, Model AR15, S/N: AR01204513)	Microbiology Laboratory Area, Room 219 Bldg 1	Instrument used for pH measurements on liquid samples.
N/A	Mettler Balance (Model PE802-S, S/N: 1128481770)	Microbiology Laboratory Area, Room 189 Bldg 1	Balances used to weight the different reagents, samples and standards during test method analysis.
N/A	Mettler Balance (Model XPR02-S, S/N: 1128501809)	Microbiology Laboratory Area, Room 219 Bldg 1	

Work Instruction	
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Title: Site Validation Program	
Doc. No. PESA-DOC-0632	Version: 15.0, CURRENT

Attachment 9: Qualified Equipment List (continuation)

Facility	Equipment	Area of Service	Description
N/A	Olympus Microscopes (2 units, Model BX-40, S/N: 7H00888 and 6E10514)	Microbiology Laboratory Area, Rooms 218 and 181 Bldg 1	Equipment used to support the Microbiology Laboratory operations.
N/A	LabLine MultiBlock Heaters (2 units, Model 2004-1BIBCE, S/N: C901822 and 12010226)	Microbiology Laboratory Area, Room 217 Bldg 1	Equipment used to support the Microbiology Laboratory operations.
N/A	Nikon Microscopes (Model: 9561, S/N: 990442)	Microbiology Laboratory Area, Room 217 Bldg 1	Equipment used to support the Microbiology Laboratory operations.
N/A	Nikon Microscope (Model: Optiphot 100, S/N: 661894)	Microbiology Laboratory Area, Room 219 Bldg 1	Equipment used to support the Microbiology Laboratory operations.
N/A	Olympus Microscope (Model: SZ, S/N: 300426)	Microbiology Laboratory Area, Room 219 Bldg 1	Equipment used to support the Microbiology Laboratory operations.
N/A	Wescor Slide Warmer (Model: 7260, S/N: 7260426)	Microbiology Laboratory Area, Room 181 Bldg 1	Equipment used to support the Microbiology Laboratory operations.
N/A	Beward Limited Stomacher (Model: 3603, S/N: 23408)	Microbiology Laboratory Area, Room 188 Bldg 1	Equipment used to support the Microbiology Laboratory operations.

Work Instruction	
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Title: Site Validation Program	
Doc. No. PSSA-DC-0382	Version: 15.0, CURRENT

Attachment 8: Qualified Equipment List (continuation)

Eqpt. No.	Equipment	Area of Service	Equipment Description
0301	REVOO Scientific Ultra Freezer (Model UL 1335-7-214, S/N: P240-037305-PG)	Analytical Laboratory Area Room 157, Building 1	Freezer with an operational range of -80 to -89 °C.
0744	Frigidaire Hotpack Refrigerator (Model 378120, S/N: C9357212)	Analytical Laboratory Area Room 157, Building 1	Refrigerator with an operational range of 2 to 8 °C.
N/A	Brookfield Viscometer (Model RVCMAH CPSS, S/N: 2149245)	Analytical Laboratory Area, Bldg 1	Instrument used for viscosity measurements of samples.
N/A	HPLC #01 (including RIUV detectors and Alliance)	HPLC Room, Laboratory Area, Room 158 Bldg 1	Instruments used as part of the Waters Millennium Data Acquisition System of the Analytical Laboratory, to acquire chromatographic data from samples.
N/A	HPLC #02 (including RIUV detectors and Alliance)		
N/A	HPLC #12 (including auto sampler, pump controller, detectors and DAD)		
N/A	HPLC #13 (including auto sampler, pump controller, detectors and DAD)		
N/A	Hewlett Packard GC's #24 (Model 6800, S/N: US11244137)	Analytical Laboratory Area, Room 158 Bldg 1	Gas Chromatography Instruments used to acquire data from samples.
N/A	Perkin Elmer FTIR (Model Paragon 1000, S/N: 45929)	Analytical Laboratory Area, Room 157 Bldg 1	Instrument used for infrared (Fourier transform) measurements on samples.
N/A	Shimadzu Spectrofluorometer (Model RF1801, S/N: A481853C0592-P)	Analytical Laboratory Area, Room 157 Bldg 1	Instrument used for fluorometric measurements on samples.
N/A	Wheeler Roller Apparatus (Model 948521, S/N: F88126)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Perkin Elmer UV/VIS (Lambda 80, S/N: 52240)	Analytical Laboratory Area, Room 157 Bldg 1	Instrument used for ultra-violet visible spectrum measurements on samples.
N/A	Mettler Toledo Balances (4 units, Model AT204, S/N: 1118502214, 1118428812, 1118516419 and 1118511811)	Analytical Laboratory Area, Rooms 157 and 158 Bldg 1	Balances used to weight the different reagents, samples and standards during test method analysis.
N/A	Mettler Toledo Balance (1 unit, Model MT-5, S/N: 1118453124)	Analytical Laboratory Area, Room 158 Bldg 1	
N/A	Mettler Toledo Balance (1 unit, Model PR 5002, S/N: 1118210021)	Analytical Laboratory Area, Room 158 Bldg 1	

Work Instruction	
Status: Effective	Effective Date: 5/5/2010 10:02:03 (EST)
Title: Site Validation Program	
Doc. No. PSGA-DOC-0892	Version: 15A, CURRENT

Attachment 8: Qualified Equipment List (continuation)

QID	Equipment	Area of Service	Description
N/A	Heraeus Oven Vantage (Model VT 6080M, S/N: 51314640)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Bernstead Thermolyse (Model F4005, S/N: 1038820161875)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Pharos Solo-400 pH Meters (5 units, Model ARTS, S/N: AFB1204800, AFB1504700, AFB1204910)	Analytical Laboratory Area, Room 157 Bldg 1	Instruments used for pH measurements on liquid samples.
N/A	Brinkman Oven (Model WTB Blister, S/N: 892200)	Analytical Laboratory Shared Area Room 154, Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Fisher Scientific Mini Centrifuge Mini Spin (Model 7209G, S/N: 58E2-07489)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Mettler Toledo Karl Fisher DL 50 (2 units, Model DL 50, S/N: 0118082438 and 5118480210)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Mettler Toledo Karl Fisher DL 50 (2 units, Model DL 50, S/N: 0115460825 and 5115460827)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Lightin Mixer (Model LabMaster 81, S/N: 803400)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Parker Briston TOC Gas Sensor (1 unit, Model TOC-1850, S/N: TOC12500240A)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Shimadzu TOC 8000/8000A (1 unit, S/N: H09104288000000)	Analytical Laboratory Area, Room 157 Bldg 1	Instruments used for TOC measurements on samples.
N/A	Spinman Melting Point Analyzer (Model E-545, S/N: 405213010002)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Perkin Elmer Polarimeter (Model 341, S/N: 7282)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Berkman Centrifuge (Model J2-MC, S/N: K05417)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
N/A	Fisher Scientific Conductivity Meters (2 units, Model AB30, S/N: AB31205503 and AB31205505)	Analytical Laboratory Area, Room 157 Bldg 1	Instrument used for the measurement of conductivity in liquid samples.
N/A	Rudolph Rectometer (Model J257, S/N: 89595)	Analytical Laboratory Area, Room 157 Bldg 1	Equipment used to support the Analytical Laboratory operations.
0580	Ampro (Star) Glass Washer (Model: Reliance 400, S/N: 862760000)	Analytical Laboratory Area, Room 155 Bldg 1	Equipment used to support the Analytical Laboratory operations.

Schedule 2
Owned Transferred IP

U.S. Patents:

Patent No.	Grant Date	Title
5427778	June 27, 1995	Gel formulations containing growth factors and acrylamide polymer
5457093	October 10, 1995	Gel formulations containing growth factors

Foreign Patents:

Patent No.	Country	Grant Date	Title
859596	EP	Dec. 5, 2001	Gel formulations containing growth factors

US Trademark:

Mark	Jurisdiction	Registration No.	Registration Date
REGRANEX	US	1928621	October 17, 1995

Foreign Trademarks:

Trademark:	Country:
REGRANEX	Australia
REGRANEX	Benelux
REGRANEX (in Korean)	Benelux
REGRANEX	Brazil
REGRANEX	Canada
REGRANEX	Ireland
REGRANEX	Israel
REGRANEX (word only)	Japan
REGRANEX	South Korea

REGRANEX (in Korean)	South Korea
REGRANEX	United Kingdom
REGRANEX	Albania (IR)
REGRANEX	Algeria (IR)
REGRANEX	Austria (IR)
REGRANEX	Belarus (IR)
REGRANEX	Bulgaria (IR)
REGRANEX	Croatia (IR)
REGRANEX	Cuba (IR)
REGRANEX	Czech Republic (IR)
REGRANEX	Egypt (IR)
REGRANEX	France (IR)
REGRANEX	Germany (IR)
REGRANEX	Hungary (IR)
REGRANEX	Italy (IR)
REGRANEX	Int'l Registration - Madrid Agreement/Protocol
REGRANEX	Kazakhstan (IR)
REGRANEX	North Korea (IR)
REGRANEX	Latvia (IR)
REGRANEX	Liberia (IR)
REGRANEX	Liechtenstein (IR)
REGRANEX	Macedonia (IR)
REGRANEX	Monaco (IR)
REGRANEX	Montenegro (IR)

REGRANEX	Morocco (IR)
REGRANEX	Poland (IR)
REGRANEX	Portugal (IR)
REGRANEX	Romania (IR)
REGRANEX	Russian Federation (IR)
REGRANEX	San Marino (IR)
REGRANEX	Serbia (IR)
REGRANEX	Serbia (old code) (IR)
REGRANEX	Slovak Republic (IR)
REGRANEX	Slovenia (IR)
REGRANEX	Spain (IR)
REGRANEX	Sudan (IR)
REGRANEX	Switzerland (IR)
REGRANEX	Ukraine (IR)
REGRANEX	Vietnam (IR)

Trademark:	Country:	Assignment Recording Date:
REGRANEX	China	Renewal due on 12/13/2016; Assignment filed 2/9/2010, recordation pending.
REGRANEX	India	Renewal due on 2/16/2011; Assignment filed 3/19/2010, recordation pending.
REGRANEX	Indonesia	Renewal due on 3/17/2015; Assignment filed 4/30/2010, recordation pending.
REGRANEX	Mexico	Renewal due on 12/14/2014; Assignment filed 5/15/2010,

		recordation pending.
REGRANEX	United Arab Emirates	Renewal due on 9/17/2016 Assignment filed (no actual date provided by local counsel), recordation pending.

Schedule 3
Telephone Line and Domain Names

Telephone Line:

1-888-REGANEX

Domain Names:

1. Regranex.be
2. Regranex.biz
3. Regranex.ca
4. Regranex.co.il
5. Regranex.com
6. Regranex.eu
7. Regranex.info
8. Regranex.mobi
9. Regranex.net
10. Regranex.org
11. Regranexcan.biz
12. Regranexcan.info

Schedule 4
Transferred Contracts

1. Manufacturing Agreement, dated March 23, 2010, between Systagenix Wound Management, Limited, and DPT Lakewood, Inc.
2. RGX Distribution Agreement (San German), dated December 1, 2008, between Systagenix Wound Management, Limited, and OMJ Pharmaceuticals, Inc.
3. Transitional Processing & Services Agreement (San German), dated December 1, 2008, between Systagenix Wound Management Manufacturing, Limited, and OMJ Pharmaceuticals, Inc.
4. Contract Management and Pricing Agreement, dated December 1, 2008, between Systagenix Wound Management (US), Inc., and Ethicon, Inc.
5. Quality and Compliance Agreement Regranex, dated November 17, 2008, between OMJ Pharmaceuticals, Inc., and Systagenix Wound Management, Limited.
6. Supply Agreement, dated July 31, 2008, between Novartis Vaccines and Diagnostics and GPSG, a unit of Ortho-McNeil-Janssen Pharmaceuticals, Inc.
 - a. Assignment of Agreements, dated July 1, 2009, among Novartis Vaccines and Diagnostics, Inc.; GPSG, a unit of Ortho-McNeil-Janssen Pharmaceuticals, Inc.; Ethicon, Inc.; and Systagenix Wound Management, Limited.
7. Master Services Agreement, dated December 1, 2008, between The Lash Group, Inc. and Systagenix Wound Management (US), Inc., together with related Statement of Work dated December 1, 2008.
8. License Agreement, dated January 18, 1994, among Novo Nordisk A/S, ZymoGenetics Inc., Johnson & Johnson, and Chiron Corporation (the "1994 License Agreement").
 - a. First Amendment to the 1994 License Agreement, dated January 1, 1997.
 - b. Second Amendment to the 1994 License Agreement, dated June 5, 2000.
 - c. Notice re: Assignment by Novo Nordisk A/S to ZymoGenetics, Inc., dated October 31, 2000.
 - d. Notice re: Assignment by Novo Nordisk A/S to ZymoGenetics, Inc., dated April 30, 2001.
 - e. Assignment of License Agreement, dated March 1, 2010, among Novartis Vaccines & Diagnostics, Inc., Novo Nordisk A/S, ZymoGenetics, Inc., ZymoGenetics, LLC, Johnson & Johnson, and Systagenix Wound Management, Limited.
9. License Agreement, dated June 10, 1986, between Chiron Corporation and Ethicon, Inc. (the "1986 License Agreement").

- a. First Amendment to 1986 License Agreement, dated August 14, 1987.
 - b. Notice re: Partial Termination of 1986 License Agreement, by Chiron Corporation to Ethicon, Inc., dated September 16, 1997.
 - c. Assignment of Agreements, dated July 1, 2009, among Novartis Vaccines and Diagnostics, Inc.; GPSG, a unit of Ortho-McNeil-Janssen Pharmaceuticals, Inc.; Ethicon, Inc.; and Systagenix Wound Management, Limited.
10. Master Agreement, dated June 10, 1986, between Chiron Corporation and Ethicon, Inc. (the "1986 Master Agreement").
- a. First Amendment to the 1986 Master Agreement, dated August 14, 1987.
 - b. Letter Agreement re: Definition, dated July 6, 1988, between Chiron Corporation and Ethicon, Inc.
 - c. Notice re: Partial Termination of 1986 License Agreement, by Chiron Corporation to Ethicon, Inc., dated September 16, 1997.
 - d. Assignment of Agreements, dated July 1, 2009, among Novartis Vaccines and Diagnostics, Inc.; GPSG, a unit of Ortho-McNeil-Janssen Pharmaceuticals, Inc.; Ethicon, Inc.; and Systagenix Wound Management, Limited.