The Variant

www.asqsefla.org

Section 1510 with Section 1515 June 2011

Inside this issue:

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Contributions will be edited to meet space requirements.

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Employment ads updated daily and are posted on the website, www.asqsefla.org

Section 1510 Website

www.asqsefla.org

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Section 1515 Website

www.asqpalmbeach.org

See All Section Activities
Certification Refresher Classes
Section Dinner Meetings
and On-Line Registrations

Message from the Section 1510 Chair By Eduardo M Suarez

Greetings to all members of Section 1510.

The topic for our last presentation was "Supplier Quality Expectations – a Panoramic 360° View" by Braulio Ortiz, co-founder and project manager of Bio Teknica located in Coral Gables, Florida. Braulio has an extensive experience in regulatory compliance, quality systems, engineering, process validation, and manufacturing for the medical device, biomedical, and pharmaceutical industries. The increased scrutiny by the FDA confirms the importance of the quality of supplied materials to medical device manufacturers and pharmaceuticals. Braulio's knowledge afforded him the opportunity to clearly explain the significance of supply chain management, purchasing controls, and that manufacturers must make sure that all aspects of the supply chain are compliant.

Our current fiscal year is coming to an end on June 30th. The last planned activity will be a tour of the Noven Pharmaceuticals, Inc facility near the Zoo Miami on June 14th. Due to security clearance, the deadline for registration is May 31st. The program starts at 12:30 with a lunch courtesy of Noven and registration. The tour is scheduled to end by 16:00. This is a great opportunity to visit a specialty pharmaceutical company engaged in the research, development, manufacture, marketing and sale of prescription pharmaceutical products.

As a reminder, The Voice of the Customer survey was sent out by e-mail by Shawn Currie and we would like everyone's input

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Message from the Section 1515 Chair by Bob Gilbert

Hello all. Section 1515 has completed our election of officers, and I am happy to announce all of our positions are filled. I would like to thank our returning Leaders and our new Chairs for accepting their positions for the 2011-2012 program year. I would especially like to thank Rae Henoch, our new Secretary, Gary Stark, our new Membership Chair, and Alex Schraff, our new Newsletter Editor, for taking on these very important positions within our Section. The Board will be meeting the second week of June to close out the old year, and begin the planning process for the new program year. We are looking for additional Committee Members to work with our Chairs to assist in implementing many of our initiatives. If you are interested in working with our Leadership Team, please contact one of our Chairs, and join us at our planning meeting.

On Thursday, May 5th, I had the pleasure of attending our annual joint meeting with ASTD at Northwood University in West Palm Beach. The presenter, Jeannine Rizzo, gave an incredible presentation to the attendees on "Aligning People and Strategy." I truly loved her real life examples, and pictures, of actual situations which demonstrated a lack of alignment within certain organizations, and how this impacts both the customer and the organizational image. Her presentation presented the benefits of establishing KFA's (Key Focus Areas), and making sure that there is alignment to them throughout the organization. KFA's are customizable, and should be established to set direction for the organization and help measure improvement. I hope to see more of our members attend these joint meetings in the future.

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Section 1510 Monthly Meeting Email Reminders!

If you would like to receive an E-Mail reminder of Section 1510 Monthly Dinner Meetings, please send a valid e-Mail address in the body of your response to Section 1510 Arrangements Chair Chair Rich Eggleston, Email: rich@pmi-inc.net. . Please place in Subject Line: "ASQ Meeting Reminder"

Section 1510 ASQ Certification Refresher Classes

To participate in an ASQ refresher class, please sign up on the web page at www.asqsefla.org. You do not need to be a member of ASQ to participate in a class. RU's are earned for class participation. If you have a question regarding registration or class site, call Mayra Bisnow, Education Chair, at 954-478-0384 or email mbisnow@gmail.com.

Please sign up at least three weeks before the scheduled class start to assure you have the textbook at first session. Any classes that do not register a minimum of five students will be cancelled. *All classes are refresher classes.*

Class	Class Name	RU's	Fee Includes materials, handouts and text where applicable	When	Time Classes are 5:30 pm to 8:30 unless otherwise listed	Instructor
IA	**Internal Auditing for Quality Systems**	1.0	$275	TBD based on demand	One day Seminar 8:30 am—5:30 pm	**Kannan Krishnan CQE, CQA, CQMa, CQIA, CSSBB, CQPA**
CQT	**Certified Quality Technician**	2.7	$400	August, January	9 sessions	**Eleanor Chilson CQE, CQMa, CQA, CQIA and Andy Vouloukos CQE, CRE**
CMQ/OE	**Manager of Quality / Organizational Excellence**	2.7	$450	August , January	9 sessions	**Kannan Krishnan**
CQE	**Certified Quality Engineer**	3.0	$450	September, March	10 sessions	**Jim Carbone, CBA, CQA, CQE, CQMa, Robert Pintavalle CQMa, CQE, CQA, CSSBB**
CSQE	**Certified Software Engineer**	2.4	$375	September, March	8 sessions	**Mercedes Massana CSQE, CQA**
CQA	**Certified Quality Auditor**	2.4	$375	September, March	8 sessions	**Jim Carbone**
CQIA	**Certified Quality Improvement Associate**	2.4	$375	September, March	8 sessions	**Kannan Krishnan**
CQI	**Quality Inspector**	2.4	$375	August, January	8 sessions	**Eleanor Chilson**
CRE	**Certified Reliability Engineer**	3.0	$350	August , January	10 sessions	**TBD**
BPR	**Blueprinting Reading**	2.4	$375	TBD	8 sessions	**Eleanor Chilson**
CCT	**Certified Calibration Technician**	2.4	$375	September, March	8 sessions	**Fred King - CCT**
CSSBB	**Six Sigma Black Belt**	3.0	$450	August, January	10 sessions	**Myriam Ochart, CSSBB & Gustavo Diaz CSSBB**
DOE	**Design of Experiments**	3.0	$400	TBD	TBD	**Robert Pintavalle**
CSSGB	**Six Sigma Green Belt**	2.7	$450	September, March	9 sessions	**Kannan Krishnan & Gustavo Diaz**
CBA	**Certified Biomedical Auditor**	2.7	$375	August, January	8 sessions	**Jim Carbone**

For Certification Exams dates and locations, please visit the ASQ Certification website at www.asq.org

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by June 10th. At our Section, your Leadership Team takes a breather during the months of July and August. Activities, meetings, presentations, etc. will start again in September.

On behalf of the Leadership Team of Section 1510, our wishes for a relaxing and enjoyable upcoming summer season.

Eduardo

Destination: Success
19th Annual Sterling Conference

The 19th Annual Sterling Conference opens on May 31st at the JW Marriott in Orlando and is the nation's benchmark for Baldrige-based best practices. Operating out of the executive office of the Governor, The Sterling Council is the official source for learning.

This year's theme**, *"Destination: Success,"*** reflects the Sterling Council's commitment to performance in all sectors including service, manufacturing, education, healthcare and the public sector.

Join us May 31 – June 3 at the JW Marriott Orlando Grande Lakes. Register now at www.floridasterling.com

- Participate in intensive pre-con sessions focused on driving performance improvement
- Learn from 50 dynamic workshops/7 comprehensive tracks with tangible take-aways you can implement when you return
- Gain knowledge from companies such as Lockheed Martin, Honeywell, Marriott, Shands Healthcare, FPL and many more
- Be inspired by three premier national keynote speakers
- Learn more about process management tools including lean, six sigma, and process mapping

The Sterling Conference is your official resource for Best Practices in business, education, healthcare, government, service, and manufacturing.

Make sure to attend the Education or Health Care Summit on Tuesday, May 31! Be one of the first to know about national, state, and local best practices.

Certification Test Sign-up Reminder

This newsletter is full of events that you can use to earn recertification points. If you need to earn your certification, here are the deadlines to sign up for the next tests that will be administered by Sections 1510 and 1515. To sign up, go to www.ASQ.org.

Aug 12 is the registration deadline for the **Oct 1** administration for the following exams: Biomedical Auditor, HACCP Auditor, Manager of Quality/Organizational Excellence, Quality Inspector, Quality Technician, Reliability Engineer, Six Sigma Black Belt.

Oct 14 is the registration deadline for the **Dec 3** administration for the following exams: Calibration Technician, Pharmaceutical GMP Professional, Quality Auditor, Quality Engineer, Quality Improvement Associate, Quality Process Analyst, Six Sigma Green-

Six Sigma Green Belt Training

Sponsored by ASQ Section 1510

Program details:

- Designed for Six Sigma enthusiasts who will perform or facilitate process improvement activities
- Suitable for all areas and all levels within an organization
- Nine Saturday sessions from 8:30 AM to 5 PM starting TBD. Classes planned to begin in March and October pending sufficient registration.
- Location: TBD
- Cost $2500 per participant (Breakfast and Lunch included) Grant money may be available to organizations through the State of Florida. For details go to www.workforceflorida.com under 'Incumbent Worker Training'.

Certification Program includes:

- Six Sigma methodology and DMAIC roadmap
- Hands on workshops on Six Sigma process improvement tools
- Hands on training on use of Minitab Software
- Successful completion and presentation of a strategically linked process improvement project.
- Six Sigma Handbook by Thomas Pyzdek and Six Sigma Memory Jogger presented as reference material

Instructors are Six Sigma Black Belt practitioners certified by ASQ:
Kannan Krishnan CSSBB, CQMa, CQE, CQA, CQIA, CQPA
Gustavo Diaz CSSBB,

Program Benefits:

- Competitively priced to accommodate individuals and company sponsored employees
- Credibility of the certifying body (ASQ Section 1510)
- Weekend program
- Included in this program (not in most other Green Belt Certification Programs): Use of Minitab Software, Design Of Experiments (DOE)
- Coverage of the 'Certified Quality Improvement Associate' (CQIA) Body of Knowledge
- EU Points for ASQ re-certification: 6.0

Prerequisites:

- Laptop preferred but **not** mandatory
- Basic math skills

Requirements for Certification:

- Attend a minimum of 6 sessions
- Present a completed project

For registration and additional information please contact section Education Chair Mayra Bisnow, at 954-478-0384 or email mbisnow@gmail.com. You may also register through our section website www.asqsefla.org

The Section 1510 Orlando Acevedo Scholarship applications are due by June 20th. More information on the 1510 website www.asqsefla.org

Congratulations to:
Joe Paparella for winning the $25 gift card at the May 10th 1510 dinner meeting door prize drawing

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Thanks to tremendous response, almost half of the seats at our upcoming joint event are sold out. With the South Florida Business Journal joining us as a sponsor, this event looks to be one of the best of its kind. Section 1515 has been working with our partners at ASTD, SME, and ACHE, to create a learning conference that will include input from all four sectors, while focusing on a common theme. The 1st Annual Joint Conference will be held at the FAU Student Union Building, in Boca Raton, on June 16, 2011. The title of the conference is "The Human Factor in Performance Improvement." The conference will focus on the human challenges to performance improvement initiatives in the fields of Quality, Healthcare, Training and Development, and Engineering. Highly recognized speakers will represent each of these specific disciplines. The Keynote Speaker, Dr. David Kopp, will synthesize all the break-out sessions in his presentation, "The Performance Formula." The six hour event will culminate with a Buffet Dinner and Networking opportunities. For more information, and to register, please log onto the ASTD website at www.ASTDSFL.org. Registration will be limited. Sign up early.

The turnout for our SME sponsored blended Lead Auditor course was great. We ended up with a waiting list of five people, and will be offer the combo course again in the fall. The course gave the participant both ISO 9001:2008 and ISO 14001:2004 Lead Auditor certifications, for the normal cost of one class. Thanks again to SME150 for working with us in offering this course. For more information about the fall course schedule, please contact gil@SME150.org.

Please check out our website for more information about our upcoming Plant Tour at the Publix Distribution Center in Deerfield Beach, on May 27th. Our final event for the program year will be the joint conference on June 16th. Please be sure to sign up soon. Our website is http://www.asqpalmbeach.org.

I hope to personally see you all at this event, and ultimately getting some new folks added to our membership and committees.

ASQ 1515 Plant Tour at Publix — May 27, 2011

"Tour of Publix Super Markets, Inc. Deerfield Beach Facility"

Publix Super Markets is ranked as one of FORTUNE's top 100 largest corporations & has been rated on their "100 Best Companies to Work For" list for 14 years straight. The Deerfield Beach facility is 1.3 million sq ft & is believed to be the largest refrigerated building in the US when built in 1988. The manufacturing area is 172,500 sq ft. The plant bottles over 1 million gallons of product per week & services 342 stores.

The presentation will include general information about Publix & the Deerfield Beach facility, Corporate Quality Assurance & the manufacturing plant Quality programs. A tour of the manufacturing operations will follow the presentation.

Date: **Friday, May 27, 2010**
Time: **2 – 4 p.m.**
Place: Publix Deerfield Beach Facility,
Cost: Free. Open to ASQ members only.

Please pre-register by email to Enrique at emb109@aol.com subject: ASQ Publix Tour
Registration closes at noon May 25. No exceptions. Capacity is limited. Please arrive about 10 min early to be able to start the program on time

Employment Opportunities

A plant in Fort Lauderdale is seeking an **Assistant Quality Manager / Laboratory Technician**. Must have **experience** in a manufacturing environment with: AIAG PPAP, ISO QMS, GDT, SPC, CMM and or Vision machine operation, CAD Software, Surface Plate Inspection Methods. **Responsibilities** include: Utilizes calibrated equipment to measure production materials to determine if they fall within acceptable parameters, Measures dimensions such as length, height, depth, angle, arc, radii, width, true position, profile and other such typical measurements on blueprints, using precision instruments such as micrometers, calipers, dial indicators, plug gauges, gauge blocks, v-blocks, sine bars, angle plates, and other such measuring equipment, Intensive monitoring of shop floor inspection process, in order to maintain a goal of zero defects.
Contact Management Representative by sending resume to quality@ebway.com

Noven Pharmaceuticals is looking for a **Sr. Manufacturing Engineer**. Responsible for the design and implementation of manufacturing processes, instrumentation and equipment from the laboratory through the pilot plant and manufacturing scale. Provides expertise engineering, design and process and/or scale-up. Assists manufacturing operation in problem solving with regards to materials, components, equipment and systems. Develops and recommends new process formulas and technologies to achieve cost effectiveness and improved product quality. Recommends processes for the production of therapeutic products using tankage, piping, coating /oven systems, laminating dies, pouching machines, and packaging systems. These pharmaceutical converting systems require automation experience with both PLC and SCADA software systems that control both intermittent or continuous motion line machines.
Requirements: BS or equivalent in Mechanical, Industrial or Chemical Engineering or a related engineering discipline with 5 – 9 years of experience and/or training; or MS with 3 – 7 years of experience and/or training; or PhD with 0 – 5 years of experience and/or training; or equivalent combination of education and experience. Experience in the pharmaceutical or medical device industries will be an asset. Must possess the basic understanding of engineering fundamentals, engineering best practices, troubleshooting methodology as well as strong familiarity of the scientific method. **Send resume to:** rkelly@noven.com

Noven Pharmaceuticals has an opening for a **Temporary Quality Assurance Associate.**. Investigate customer complaints for Noven products in the TrackWise System and perform all activities related to investigation of a complaint, including processing return samples for investigation or credit and requesting data from other departments. Assure on-time closure of complaint records and associated TrackWise tasks. Monitor complaint activity for trends and initiate trend investigations as appropriate. Identify critical complaints, notify supervisor and prioritize investigations to met reporting requirements. Provide weekly progress reports and metrics. Participate in team activities and projects as assigned.
Requirements: Candidate must have a science-related degree with at least 3 years experience in the Pharmaceutical or Medical device industry. Working knowledge and familiarity with the process for investigation of product quality complaints in the TrackWise software system is a plus. The candidate must have excellent written and verbal communication skills and ability to manage multiple tasks at a time. Should be ability to work independently or in a team environment, as needed. **Send resume to:** rkelly@noven.com

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1510 Plant Tour

Plant Tour at Noven Pharmaceuticals Inc.,
Location: 11960 SW 144th Street Miami, FL 33186 Tel: 305-253-5099
Date & Time: Tuesday June 14th, 2011 12:30 PM-16:00 PM starting with LUNCH.

Noven Pharmaceuticals Inc., is a specialty pharmaceutical company engaged in the research, development, manufacture, marketing and sale of prescription pharmaceutical products. Noven has over 650 employees working in facilities located in: Miami, FL; New York, NY; and Carlsbad, CA, and in sales force territories located across the U.S.
Noven has offered ASQ a Plant Tour to demonstrate their engagement in the field of Quality.
Dr. Barry Gujral, the Director of Quality Engineering at Noven, will be our host during the Plant Tour.

The Plant Tour starts at 12:30 PM with **Lunch and Registration**.
We will be visiting the production Line followed by a presentation by the Management of Noven.

You are invited to attend this Plant Tour and we are excited to meet you there on June 14th.
Due to Security Clearance requested by Noven, the **DEADLINE for registration is May 31st, 2011.**
Please go to www.asqsefla.org to register. For information about the Plant Tour, please contact Saeid Ebrahim our Program chair by Email: sebrahim@bioteknica.com or Cell: 954-610-4792

Let us know what you think

Section 1510 has sent an e-mial to our section members to participate in our 1st Voice of the Customer Survey. This survey is intended to get input on areas in which we can improve our section in order to provide the best experience for our members. You have received an e-mail in from Shawn Currie that will have a link to the survey. There is also a link on www.asqsefla.org. It is approximately 20 questions, and should take 10-15 minutes to complete. Please respond by June 10. We hope to have 100% participation. This is your opportunity to have your voice heard (anonymously).

Virtual Refresher Course Available —- Manager of Quality/Organizational Excellence Certification - CMQ/OE

The Certified Manager of Quality/Organizational Excellence is a professional who leads and champions process-improvement initiatives—everywhere from small businesses to multinational corporations—that can have regional or global focus in a variety of service and industrial settings.
A Certified Manager of Quality/Organizational Excellence facilitates and leads team efforts to establish and monitor customer/supplier relations, supports strategic planning and deployment initiatives, and helps develop measurement systems to determine organizational improvement.
The Certified Manager of Quality/Organizational Excellence should be able to motivate and evaluate staff, manage projects and human resources, analyze financial situations, determine and evaluate risk and employ knowledge management tools and techniques in resolving organizational challenges.
If you are interested in becoming a Certified Manager of Quality/Organizational Excellence, the next exam date is **Saturday, October 1, 2011**. To help you pass the exam on the first try, ASQ Olde Colony is once again offering our successful virtual **Certified Manager of Quality/ Organizational Excellence Refresher Course, starting Monday July 18th**.
Our experienced instructor, **Anthony DeMarinis,** is a Certified Manager of Quality/Organizational Excellence and a Certified Quality Auditor-Biomedical with over 15 years experience using Quality Management techniques and value added auditing, including direct involvement with FDA and ISO audits.
For registration or more information, please contact Rochelle Jones at (508) 577- 0788 or education@asqoldecolony.org

Employment Opportunities

Noven Pharmaceuticals is looking for a Supervisor, **Quality Control Raw Materials**. **Responsibilities**: Develop and implement methodologies for the qualification and testing of raw materials, in-process, and components utilized in commercial and developmental finished products; troubleshoot analytical instrumentation within the laboratory. Supervise all the activities related to raw material, in-process, and component testing. **Requirements:** Bachelor's degree in chemistry. Five or more years of professional experience as well as broad knowledge of scientific principles, chromatographic analysis, analytical methods (i.e., HPLC, UPLC, GC, UV), and excellent understanding of separation theories, all types of compendia testing (USP, EP, BP, JP) and wet chemistry techniques is required. Must have excellent English written and verbal communication skills; must be highly organized and be able to work under pressure; must have good interpersonal and leadership skills; must be computer literate with knowledge of Microsoft business office applications. A background in scientific software is required. Knowledge of Water's Corp. Empower and LIMS as well as TrackWise is a plus. **Send resume to:** rkelly@noven.com

Quality Manager — Boca Raton area medical device manufacturer requires a responsible professional with FDA, QC-RA management experience who is a self-starter and a person who pays attention to details. **Education/ Experience:** Must have professional experience as well as a broad knowledge and an excellent understanding of FDA device compliance and ISO 9000 and ISO 13485 Systems. Must have a related degree with at least five years of experience in medical device industry. **Responsibilities:** Ensuring that quality system requirements are effectively established and effectively maintained; and reporting on the performance of the quality system to management with executive responsibility for review, Performing: Control of documentation, corrective and preventive actions, disposition of nonconforming product, and coordinating customer complaint investigations, approving or rejecting all procedures or specifications, approve or reject all components,
Send resume including references to: shirley.knaster@yahoo.com

SAVE THE DATE AND REGISTER TODAY FOR A POWERFUL ANNUAL INTERDISCIPLINARY EVENT

June 16, 2011
"The Human Factor in Performance Improvement"

To be held at Florida Atlantic University, Boca Raton Student Union Building

A Joint Conference Presented by the Local Chapters of:

ASQ (American Society for Quality)
SME (Society of Manufacturing Engineers)
ACHE (American College of Healthcare Executives)
ASTD (American Society for Training and Development)

Keynote Speaker: **Dr. David Kopp** will deliver **"The Performance Formula".**
How workplace performance is the result of Knowledge, Skills, Attitudes, Motivation, and Environment.

Schedule:

1:00 pm **Registration and Networking**
1:45 pm **Opening and Welcoming Remarks**
2:30 to 3:30 pm - First Concurrent Breakout Session
Doron Zilbershtein (ASQ) - "The Hidden Aspects of Performance Improvement: The Psychology Behind the Process."
Gil Lugo and Bob Gilbert (SME) - "Battle of the Improvement Methodologies: The Human Side."
3:45 to 4:45 pm - Second Concurrent Breakout Session
Michael Sabbag (ASTD) - "The 7 Levers of Exemplary Performance"
Roger Chen, FACHE (ACHE)-"Developing Leaders to Create a Continuous Improvement Culture"
5:00 to 5:45 pm: **Keynote Speaker Dr. David Kopp**
5:45 to 7:00 pm: **Dinner and Networking**

REGISTRATION*

$75 - Early Bird Registration (before April 30, 2011)
$95 - Registration (after April 30, 2011)
Your conference registration fee includes:
- Access to all Sessions
- CE credits (as applicable)
- Refreshments
- Dinner and Networking

For More Info & to Register **www.ASTDSFL.org**
*Space is limited so register early

ASQ Re-certifications
Congratulations from the
Section 1515 Lead Team!

James Altmann
CQA, CQE

SECTION 1515 LEADERSHIP TEAM 2011-2012

Position	MemberName	Email
Chair	Bob Gilbert	bob@GreenAndSustainableSolutions.com
Arrangements Chair	Mikhail Bunich	mbunich@mapei.com
Secretary	Rae Henoch	rae.henoch.ctr@autec.navy.mil
Treasurer	Jeremy Montanti	jeremy@dmcconline.org
Auditing Chair	Charlotte Williamson	charlotte.williamson.ctr@autec.navy.mil
Nominations Chair	Jay Stahan	jstahan@qmedcorp.com
Certification Chair	Judith Gottesman	judy_gottesman@ncci.com
Recertification Chair	Judith Gottesman	judy_gottesman@ncci.com
Program Chair	Gil Lugo	gil@GreenAndSustainableSolutions.com
Membership Chair	Gary E Stark	gary.stark.ctr@autec.navy.mil
Education Chair	Gil Lugo	gil@GreenAndSustainableSolutions.com
Internet Chair	Mauricio Perry	mauricio.perry@cslplasma.com
Publicity Chair	Denise Sandberg	sandimountain@aol.com
Newsletter Editor	Alexander Schraff	aschraff@eeellc.net
VOC Chair	J ay Stahan	jstahan@qmedcorp.com

Employment Opportunities con't

Ads for individuals seeking positions are free to ASQ Section 1510 / 1515 members and subject to space availability.
Ads for open positions in a quality related field are free and not limited to ASQ members.
To place an ad, contact Placement Chair, Eleanor Chilson, (954)661-5197, or chilsone@bellsouth.net

Verify / VSC is continually searching for **Independent Contractors** with a background in Quality. Experience in the Defense and Aerospace Industry preferred. If you are interested, please **send all inquiries via email to recruitment@vscnet.com.** You can also visit our website at www.vscnet.com for more information on our company.

QA professionals having backgrounds in engineering, manufacturing and quality. Gaumard designs and manufactures more than 200 simulators for health care education. These products include mechanical, electronic and software components. Responsibilities include developing procedures for inspection and testing of incoming components, work in process, and finished goods; reporting quality and reliability data; evaluating and reporting upon customer returns. **Send resumes to john@gaumard.com.**

BE Aerospace in Medley, FL has an immediate opening for a **Quality Engineer, Responsibilities:** Performs professional Quality Engineering assignments of considerable complexity under direction with considerable discretion as to work details, Designs, installs and continually evaluates complex quality assurance and/or control methods and systems, Designs fixtures and processes to insure the continuing maintenance of product quality; may conduct vendor visits to insure quality; analyzes reports and returned products to determine quality trends and recommends correction action., May participate in materials review activities to determine material disposition., Develops standards and procedures to provide quality guidance and methods. **Requirements:** .Bachelor's degree in Engineering from four-year college or university; and minimum three years related experience and/or training; or equivalent combination of education and experience. Ability to read, analyze and interpret general business periodicals, professional journals, technical procedures or governmental regulations. Ability to write reports, business correspondence and procedure manuals. Ability to effectively present information and respond to questions from groups of managers, clients, customers and the general public. Please visit the following website to apply:
https://beaerospace.tms.hrdepartment.com/cgi-bin/a/highlightjob.cgi?jobid=8216&lcid=en-US&is_preview=1

BE Aerospace also has openings for a **Senior Customer Quality Engineer, QA Inspector II** and a **QA ODA Inspector III.** If interested in any of these positions, please contact: **tetiana_danylchenko@beaerospace.com**. Details for each of these positions are below.
Senior Customer Quality Engineer: Responsible for product excellence and implementation of quality assurance programs to provide defect free products to all customers. JOB REQUIREMENTS: BS/BA degree in an applicable field (engineering) with a minimum 5-7 years of Quality Engineering experience, Strong Problem Solving skills with over 4 years writing Corrective Actions, Excellent English communication skills, both verbal and written, ,Must be advanced in MS Word and Excel and PowerPoint, Must have Leadership skills leading teams, Must have over 3 years working with Customers, Must have Aerospace Industry Knowledge, Working knowledge of AS-9100/9120 Quality standard, ADDITIONAL SKILL: Knowledge of Lean Principles, Professional quality certifications (ASQ-CQE,CQA), Six Sigma Greenbelt certification, Strong knowledge of FAA regulations
Quality Assurance Inspector II: Inspects incoming parts/material, verify that certification is correct, checks parts for processing, check special orders, in charge of micrometers, gauge controls, calipers, indicators, etc. Provides leadership to the Inspection team. JOB REQUIREMENTS:
Minimum of 4 years of experience in QA inspection methods and procedures, Fluent in English, reading, writing and verbal communication, Strong analytical, organizational and troubleshooting skills, Required to pass documented color and near vision exam, Knowledge of AS-9100 Requirements, Strong inspection knowledge with use of Calipers, ring/pin gages, comparator and smart scope, High school education or GED equivalent, ADDITIONAL SKILLS: Associate degree, Understanding of Six Sigma / team concepts preferred, ISO knowledge and understanding preferred.
Quality Assurance ODA Inspector III: Works on behalf of the FAA Organizational Designation Authorization (ODA) program as a Unit Member (UM). This individual will visually and dimensionally inspect aerospace product and documentation to ensure conformity to Type Design and safe operation, issue FAA airworthiness certification and generate all paperwork to ensure compliance with any and all FAA export requirements.
JOB REQUIREMENTS: Minimum 5 years experience with inspecting Aerospace products of similar complexity. (Ie..Nuts, bolts, o-rings, bearings, springs, Hi-Loks, terminals, gaskets, decals, wires, pins, assembles parts, brackets, screws, rivets and electrical components), Must be able to work effectively with the FAA and the ODA Site Lead, Certified ODA Unit member(The selected person must be capable of obtaining FAA approval as a Honeywell ODA Unit member within 6 months of hire date), Proficient at reading and interpreting Type design data (i.e. Blue Prints, specifications, Standards (AN,MS,NAS,NASM, MIL) and FAA related documents/ regulations), Extensive knowledge in the use of inspection measuring equipment such as Calipers, Micrometer, Optical Comparator, Johnson gages, ring and pin gages, plug gages, block gages and microscope, Proficient at interpreting and understanding hardware manufactures certification documents such as Manufactures Certificate of Conformance, Chemical/ Physical Test, special process certification, independent lab reports and First Articles reports, Knowledge of AS-9100/ 9120 Quality Standards, Must demonstrate good Leadership skills, High school diploma or equivalent, ADDITIONAL SKILLS: Bachelor Degree in related field, Experience dealing with regulatory agencies/requirements preferred, Previous DMIR, DAR, ODA unit member experience preferred, Prior or current Airworthiness Designee certification (ODAR,ODA DAR, DMIR) is preferred. Proficient computer skills

Seeking an opportunity to continue a career path in Quality Management Systems: Experienced Quality Manager of a **Plastic Extrusion Molding** company and a **Certified Lead Auditor for ISO-9001:2008 QMS**. Responsible for all Quality Management Systems per ISO-9001:2008 requirements, preparing quality assurance documentation and systems conceptual models, interacting with quality system registrars, managing customer quality interactions, and overseeing supplier development interactions. Primary products service the medical industry. Familiar with ISO-13485 and FDA regulatory requirements. **Contact:** Joseph E. Satek, jsatek03@yahoo.com Cell: 412-780-5087

A small biotech location in the Southern Caribbean is looking for a Director of Quality Systems. The parent company is here in the US. The client wants someone who has already lived on a Caribbean island before. They process plant materials through fermentation, sterilization, and lyophilization. They would like at least 15 years of experience to include some microbiology responsibility. This is a permanent, full time position. They provide an excellent relocation package, very good salary, bonus, including foreign service pay and housing allowances. See 1510 website for full job description. **Contact**: Dr Thomas H. Agrait, I.E.Lean Six Sigma Master, President - CEO, Lean Enterprise Consulting,Inc, USA - Latin America - Europe - India - Asia, Tel 954-258-3117, tagrait@gmail.com

Please visit the Career area of the Section 1510 and 1515 websites for the most current and complete listing of opportunities

ASQ AMERICAN SOCIETY FOR QUALITY

The Variant

Section Management Process TOTAL QUALITY AWARD 2006-07

Section 1510 with Section 1515 *June 2011*

Plant Tour at Noven Pharmaceuticals Inc.,

Location: 11960 SW 144th Street Miami, FL 33186 Tel: 305-253-5099
Date & Time: Tuesday June 14th, 2011 12:30 PM-16:00 PM starting with LUNCH

Noven has offered ASQ a Plant Tour to demonstrate their engagement in the field of Quality.
Dr. Barry Gujral, the Director of Quality Engineering at Noven, will be our host during the Plant Tour.

Noven Pharmaceuticals Inc., is a specialty pharmaceutical company engaged in the research, development, manufacture, marketing and sale of prescription pharmaceutical products.

Due to Security Clearance requested by Noven, the DEADLINE for registration is May 31st.
Please go to www.asqsefla.org to register.
For information about the Plant Tour, please contact Saeid Ebrahim our Program chair by Email: sebrahim@bioteknica.com or Cell: 954-610-4792