M/F SENIOR PRE-CLINICAL SCIENTIST

- **About us**

Advanced Accelerator Applications (AAA) is an innovative radiopharmaceutical company that develops, produces and commercializes Molecular Nuclear Medicine (MNM) products. AAA's lead therapeutic product candidate, Lutathera®, is a novel MNM compound that AAA is currently developing for the treatment of neuroendocrine tumors, a significant unmet medical need. Founded in 2002, AAA has its headquarters in Saint-Genis-Pouilly, France. AAA currently has 22 production and R&D facilities able to manufacture both diagnostics and therapeutic MNM products, and has over 500 employees in 13 countries (France, Italy, UK, Germany, Switzerland, Spain, Poland, Portugal, The Netherlands, Belgium, Israel, U.S. and Canada). AAA reported sales of €88.6 million in 2015 (+27% vs. 2014) and sales of €81.3 million for the first 9 months of 2016 (+23% vs. 9 months 2015). AAA is listed on the Nasdaq Global Select Market under the ticker "AAAP". For more information, please visit: www.adacap.com.

AAA has a broad pipeline of products in development. In addition to the company lead product, Lutathera®, and its Gallium-68 Dotatate product NetSpot recently approved by the FDA, Phase II studies have been initiated with Annexin V-128, a diagnostic 99mTc tracer for apoptosis and necrosis, and the company has other imaging and therapeutic radiopharmaceutical products in development.

AAA is looking for a **Senior Pre-Clinical Scientist** to be based in Ivrea (To), Italy. This position will report to the Manager of Pre-Clinical Development.

- **Role**

The selected candidate will be in charge of the overall operational management and oversight of pre-clinical studies. He/She will work closely with all R&D members as well as Regulatory Affairs team and Project Management team to coordinate and ensure the successful completion of the pre-clinical activities within the agreed timelines and budget, and in compliance with the required quality and Good Laboratory Practice and ICH Guidelines.

- **Key tasks**

- Coordinate activities of pre-clinical development projects, monitor and document progresses and escalate issues to management.
- Collaborate in the design, review and approval of pre-clinical studies and study documents including but not limited to study protocols and amendments, investigator brochures, progress reports, final pre-clinical study reports, presentations.
- Prepare/ review pre-clinical documentation and ensure on time availability of critical documents and progress reports.
- Prepare pre-clinical project plan and monitor related budgets.

- Be accountable for the deliverables of pre-clinical activities within the agreed timelines and budget.
- Organize, either directly (in house) or through selected CRO that study activities are aligned to the established planning, timelines and budget.
- Ensure the appropriate communication among all functions within the company and with external vendors.
- Coordinate pre-clinical documentation and submissions in support of clinical studies and submissions to Regulatory Agencies within appropriate project timelines, in collaboration with the AAA Regulatory department.

- **Professional skills & experience**

- Advanced scientific degree with emphasis on pre-clinical development.
- Thorough knowledge of pre-clinical research and development processes.
- Comprehensive and practical knowledge of the principles of Good Laboratory Practices and the conduct of pre-clinical activities in accordance with ICH and other preclinical-related regulations/guidelines.
- 5 to 10-year pre-clinical research and development experience including in-vitro and in vivo activities.
- Experience in *in-vitro* and *in-vivo* pharmacology, PK/PD and toxicology is required.
- Experience in research and development in the field of nuclear medicine field is a plus.
- Professional proficiency in both spoken and written English

- **Personal skills & abilities**

- Strong organizational and time management skills
- Highly proactive, self-motivated, professional and dedicated
- Flexibility and ability to prioritize and manage multiple tasks in a challenging environment
- Excellent English written and oral communication skills; other languages an asset
- Proficient PC skills: Word, Excel, PowerPoint, email.
- Quality oriented with attention to details
- Accept to travel as required

- **Submission of your application:**

Please send your application in English via email to HRItaly@adacap.com. Your application shall include all the information you consider relevant, and at a minimum, a motivation letter, your personal data, education, employment history and details on your current position and salary as well as your salary expectations for this position.

Please note that we will only reply to candidates that we wish to bring to interview stage.

For further information on the company, please visit our web site www.adacap.com