

<b>Job Title:</b>	<b>SAS Programmer / Statistical Programmer</b>		
<b>Reports To:</b>	<b>Scientific Data Manager</b>	<b>Grade:</b>	
<b>Function:</b>	<b>Scientific Product Stewardship</b>	<b>Location:</b>	<b>Southampton</b>
<b>Purpose:</b>	<b>Provide programming support across R&amp;D projects. Facilitate SAS programming and Statistical programming to ensure the creation of datasets for regulatory reporting.</b>		
<b>Dimensions:</b> <small>(complete/delete/add dimensions as appropriate)</small>	<b>Staff: No direct reports.</b> <b>Other: The job holder will need to work across groups within GR&amp;D, facilitating SAS programming activities for a broad range of GR&amp;D projects to be used for regulatory submissions.</b>		

<p><b>Principal Accountabilities</b></p> <p><b>Operational/Professional/Business</b></p> <p>Apply SAS programming and analytical skills using BASE/SAS, SAS/STAT, SAS/GRAPH and SAS MACROS</p> <p>Apply SAS programming to create datasets according to CDISC standards and internally developed data standards.</p> <p>Design, develop, evaluate, validate and modify computer programs using SAS to analyse pre-clinical, clinical and survey data.</p> <p><b>Management</b></p> <p>Application of formal project management.</p> <p>The job holder will need to contribute to development and maintenance of standards for programming activities oriented to regulatory submissions and ensuring that best data management practices are followed.</p> <p>Budget. The role holder is not a budget owner, however, will be expected to contribute to the planning of budgets to manage expenditure and monitor output from the external contracts and collaborations.</p> <p><b>Leadership</b></p> <p>Role holder will be expected to provide appropriate support &amp; coaching to other team members in SAS programming.</p> <p><b>Relationship</b></p> <p>Interact with other key stakeholders to ensure timely deliverable of data used for regulatory submission.</p> <p>Participation in industry forums &amp; working groups</p> <p><b>Innovation</b></p> <p>Decisions are made based on data interpretation gained from extending what is already known in the technical area.</p>
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<b>Knowledge, Skills and Experience</b>		
<b>Qualification</b>	Directly relevant professional experience demonstrating advance SAS programming skills and SAS statistical programming	
<b>Institution Membership</b>	NA	
<b>Competencies / Experience</b>	Please list what experience the person would need to do the job (not Years of Experience)	
	<b>Essential</b>	<b>Desired</b>
Technical capability and knowledge	<p>Bachelor's degree in Computer Science, Statistics, Mathematics, otherscientific subject</p> <p>pharmaceutical/biotech or CRO setting using SAS with clinical and/or non-clinical experience</p> <p>Able to create offline listings</p> <p>Excellent working knowledge on CDISC standards with emphasis on CDASH, STDM and ADAM</p> <p>Able to develop appropriate SAS macros to facilitate data and programming checks</p> <p>Experience creating Define.xml files as per CDISC standards</p>	
Methods / Processes	Understanding of statistical methods and application of Data Standards to clinical and non-clinical studies.	
Techniques	Highly proficient in Statistical SAS programming. R programming experience a plus	
Product Knowledge	Product knowledge is not essential	Familiarity with categories of new emerging nicotine products
Business Acumen / Commercial Knowledge	Maintains up to date of new emerging products / brands are sold globally	
Leadership / Managerial Skill	Proactive; self-motivated individual with ability to work in team environment. Good communication/interpersonal skills	
Behaviours	Interacts with internal stakeholders and cross-functional teams to ensure project delivery.	