

## Project proposal template – Faculty studentships Summer 2014

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<i>Project title</i>	<input style="width: 100%; height: 100%;" type="text" value="The covalent attachment of bio-compatible polymers to proteins to improve their pharmaceutical"/>	<i>Director of Study</i>	<input style="width: 100%; height: 100%;" type="text" value="John Fletcher"/>
<i>Second Supervisor</i>	<input style="width: 100%; height: 100%;" type="text" value="Alex Sinclair"/>	<i>School</i>	<input style="width: 100%; height: 100%;" type="text" value="Pharmacy and Chem"/>
<i>Other members of supervisory team</i>	<input style="width: 100%; height: 100%;" type="text" value="Gianpiero Calabrese"/>	<i>Any requirements from applicant (eg degree in specific subject area)</i>	<input style="width: 100%; height: 100%;" type="text"/>
<b>Project summary (max 1,000 characters)</b>			
<p>Protein PEGylation, the covalent attachment of polyethylene glycol (PEG) to a protein is often used to increase the circulatory half-life by increasing the protein's hydrodynamic size whilst reducing immunogenicity and antigenicity. Protein PEGylation often results reduced toxicity and dosing frequency, and is cost effective and cost saving.</p> <p>Proteins are often formulated as lyophilised powders for reconstitution rather than solutions to maximise their stability. Unfortunately PEG has a low glass transition temperature which leads to an increased amount of molecular motion in the dried state. This can result in a loss of protein activity during lyophilisation of PEGylated proteins.</p> <p>The aim of this PhD project will be to develop methods to circumvent this problem by covalently attaching common pharmaceutical proteins such as G-CSF or interleukin, to a range of bio-compatible polymers through click chemistry and subsequent screening for improved drug delivery properties.</p>			