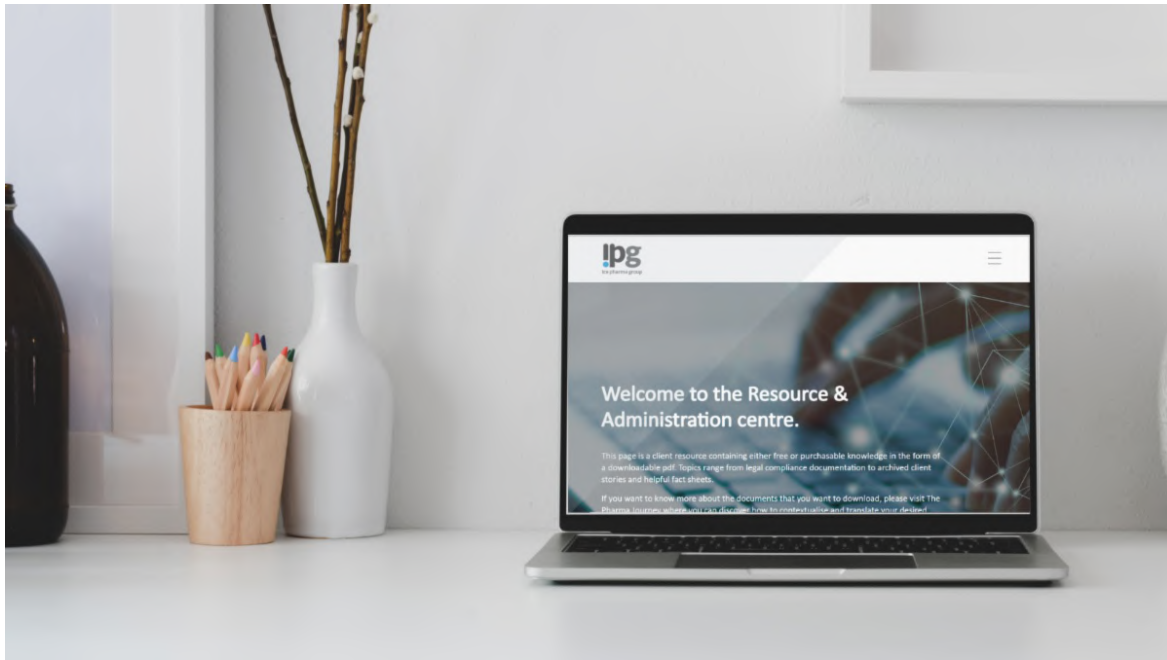


New feature coming soon...



Introducing our new feature: **Compliance information documents for purchase.**

Ahead of time we are announcing an exciting new development, the opportunity for our customers to download our compliance instruction documents directly from our website for a fixed price.

These invaluable documents include specifics on the responsibilities of WDA, MIA and MAH holders. We will also provide documentation on API licenses, GDP, GMP, QMS systems and more.

If this is something you are interested in learning more about, or if you are seeking the knowledge and tools to navigate pharmaceutical compliance to grow your business, watch this space...

[Enquire here for more information](#)

For a taste of what is to come, download our compliance controls summary document for free below.



This summary document outlines all you need to do in order to achieve compliance when handling pharmaceutical products in Europe and the UK.

This document explores a range of compliance necessities from licenses to audits to personnel and more.

The information and implementation presented within this summary may seem overwhelming.

That's where we come in.

We also offer complementary service solutions to help you achieve the compliance milestones outlined within this summary, [inquire today](#).

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