#### **Publications**

#### Guidance documents

- Guidelines for Phase I clinical trials
- First in human studies study placement, design and conduct
- Demonstrating value with Real World Data
- Guidance notes on the management of adverse events and product complaints from digital media
- Guidance notes on collection of safety information by employees and agents of pharmaceutical companies
- Guidance notes on the collection of adverse events and product quality complaints from market research programmes
- Guidance notes on UK data protection in post-marketing pharmacovigilance

#### **Industry** information

- The changing shape of academic collaborations with the pharmaceutical industry
- An insight into careers for doctors with the UK pharmaceutical industry
- Poster highlighting careers in the pharmaceutical industry
- Animals and medicines research

#### White papers

- The stratification of disease for personalised medicines
- The vision for Real World Data harnessing the opportunities in the UK
- Pharmacovigilance and the internet: a call for change
- Biosimilars position paper

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#### To learn more visit

Research, Medical & Innovation on the ABPI website: www.abpi.org.uk

ABPI member zone:

https://extranet.abpi.org.uk

ABPI schools site:

http://www.abpischools.org.uk

Careers in the pharmaceutical industry site:

http://careers.abpi.org.uk

ABPI exam site:

https://exams.abpi.org.uk

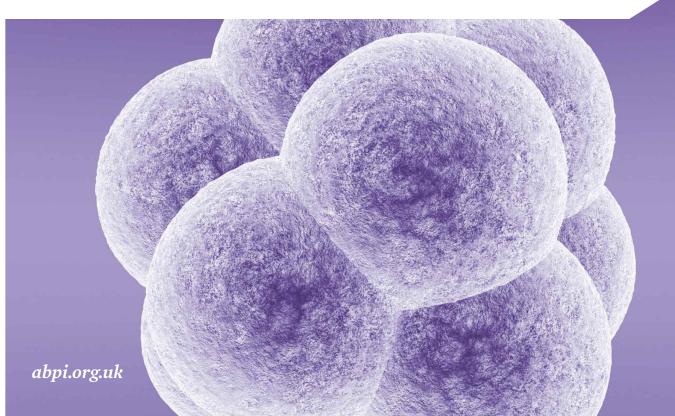
#### Contact us

abpi@abpi.org.uk



# Research, Medical & Innovation

- Creating a leading environment for R&D
- Maximising the impact of industry's investment in R&D



### We cover

Discovery

Nonclinical research

Open innovation

Education and skills

Pharmaceutical industry careers

Personalised medicines

Experimental medicine

Late phase research

Clinical trial monitoring

Medical affairs

Information about clinical trials (transparency)

Pharmacovigilance

Regulatory affairs

Manufacturing and product quality

Real World Data and patient outcomes

Environment and health & safety
eHealth and health informatics
Pharmaceutical security
ABPI Medical Representative Exam

## Key outputs in partnership

All ABPI recommendations incorporated into the Government's Strategy for UK Life Sciences

Cross-industry funded collaborations in drug safety and the 3Rs (replacing, reducing and refining the use of animals in research):

- ABPI Centre for Drug Safety Sciences collaboration
- ABPI NC3Rs collaboration

Effective transposition of the EU laboratory animal directive (2010/63/EU) enabling biomedical research, raising animal welfare standards and ensuring a level playing field across Europe

Creation of the Concordat for Openness on Animal Research

Introduction of UK patent box and improvement of R&D tax credits fostering inward investment in R&D and manufacturing in the UK

Leading thinking in pharmacovigilance including suite of best practice guidelines

More than 80 case studies on our Careers website





Open Innovation initiatives driving industry/academia collaborations eg

- MRC-ABPI Inflammation and Immunology Initiative
- MRC-ABPI Diabetes Initiative
- Translational research partnerships

Working with other stakeholders to ensure effective transposition and implementation of the EU Falsified Medicines Directive (2011/62/EU)

Over 60,000 visitors per month to the ABPI's website for schools

Shaping the UK stratified medicine agenda eg with the Technology Strategy Board's innovation platform

Leading thinking on key industry issues, including pharmaceuticals in the environment, biosimilars, regenerative medicine, early access to medicines and data protection

Development with the Carbon Trust of a computer-based tool to aid in carbon footprinting of medicinal products

Joint working to develop the ABPI *Real World Data Guidance* and subsequent training for industry