

Date	Title
30/03/2017	Guest blog: New Scottish SPIRE campaign
06/04/2017	Launch of Understanding Patient Data website
06/04/2017	New words and pictures to explain anonymisation
12/04/2017	Launch of UPD
28/04/2017	Caldicott Review Update
15/05/2017	UPD response to NHS cyber attack
15/05/2017	UPD response to concerns about Royal Free London and DeepMind deal
19/05/2017	Horizon scanning and new technologies using patient data
26/06/2017	What do we mean by 'public benefit'?
03/07/2017	ICO ruling on Royal Free London and DeepMind deal
05/07/2017	DeepMind Health Independent Review Panel first annual report
10/07/2017	New technologies using patient data - meeting summary
12/07/2017	Open letter from charities following the Government's response to the National Data Guardian Review
12/07/2017	Government response to the Caldicott Review
17/07/2017	Guest blog: data management and use - governance for the 21st century
24/07/2017	Guest blog: unifying the medical research charities conversation on data
18/09/2017	Guest blog: how giving patients access to health records can transform care - a patient's perspective
18/09/2017	Guest blog: patient data saves lives, let's acknowledge it
08/12/2017	Guest blog: who has access to data about my treatment and care?
14/12/2017	Guest blog: how my work with GPs uses patient data to improve patient safety
15/12/2017	Guest blog: how giving patients access to health records can transform care - today and in the future
05/01/2018	A look ahead: patient data in 2018
15/01/2018	UPD response to release of cancer patient data to tobacco company
16/01/2018	Memorandum of Understanding on data-sharing between NHS Digital and Home Office
07/02/2018	Cybersecurity in the NHS
08/02/2018	Guest blog: why the MoU is not in the public's interest - BMA's views
05/03/2018	Guest blog: how giving patients access to health records can transform care - a GP's perspective
06/03/2018	An introduction to the GDPR - and what it means for patient data
09/03/2018	Update on the Home Office - NHS Digital Memorandum of Understanding
12/03/2018	Data saves lives: animations
16/04/2018	Health Select Committee repeats call to suspend data sharing for immigration purposes
19/04/2018	Guest blog: voluntary sector joins the conversation on patient data
23/04/2018	Guest blog: understanding the NHS isn't easy
26/04/2018	Data for public benefit: balancing the risks and benefits of data sharing
11/05/2018	Changes to the Home Office - NHS Digital Memorandum of Understanding
18/05/2018	Guest blog: New Healthwatch England survey - careless data handling costs lives
23/05/2018	Local health and care record exemplars announced
25/05/2018	Why an opt-out rather than an opt-in or consent?
25/05/2018	Is the new national data opt-out just care.data all over again?
25/05/2018	What's the potential impact of the national data opt-out?
11/06/2018	Charities explain how patient data underpins health research
15/06/2018	DeepMind Health Independent Review Panel second annual report
28/06/2018	Future uses of data and technology in the NHS
03/07/2018	TPP error for "Type 2" opt-outs
31/07/2018	Guest blog: how the voluntary sector has helped shape the national data opt-out
21/08/2018	Guest blog: what does the public find reasonable when it comes to uses of patient data?

31/08/2018 Guest blog: patient data at the Edinburgh fringe festival

12/09/2018 Response to the consultation on the Centre for Data Ethics and Innovation

02/10/2018 Guest blog: how can third sector organisations support a transformation of patient care through data?

19/10/2018 Summary report - healthcare data: how do we get it right?

29/11/2018 Principles that put people at the heart of new data technologies in health

05/12/2018 Letter to the Secretary of State

10/12/2018 Reflections on Understanding Patient Data

08/01/2019 Next steps for Understanding Patient Data

09/01/2019 Joining up work and busting jargon: an update from our December event

15/02/2019 Public engagement in data sharing projects

11/03/2019 Listening to patients and publics: a mix of methods, one coherent approach

26/03/2019 Response to the National Data Guardian's consultation on priorities

09/05/2019 Guest blog: people powered genomics

13/05/2019 Public engagement on partnership models involving NHS data

16/05/2019 Guest blog: using National Clinical Audit data to improve care quality

28/05/2019 Introducing the UPD resource quick-guide

17/06/2019 Transparency: what is it and how do we improve it?

17/06/2019 Is being informed about patient data enough?

30/07/2019 Guest blog: building public trust in the use of health information to improve health and care

05/12/2019 Hello again

10/12/2019 Trading patient data: our view

13/12/2019 Safeguarding health data: 3 policy asks of the new Government

20/01/2020 We're hiring! Strategy and Engagement Manager role

13/02/2020 Using NHS Digital data to improve care

25/02/2020 We're hiring! Partnerships and Community Manager role

28/02/2020 Guest blog: how can patient data help us understand complex rare diseases?

02/03/2020 Accountability, transparency and public participation must be established for third-party use of NHS data

19/03/2020 Covid-19: we're here for our community

17/04/2020 Health data and Covid-19: taking action while maintaining trust

21/04/2020 Easier said than done: the challenge of transparency during Covid-19

28/04/2020 How to talk about data without talking about data

29/05/2020 The issue with tissue: towards a national registry of human tissue sampling

16/06/2020 Data in the time of Covid-19

30/06/2020 Creating the right conditions to involve the public in policymaking

08/07/2020 A new approach to decisions about data

28/07/2020 Putting the trust in Trusted Research Environments

04/08/2020 There is no single truth about Covid-19

02/09/2020 What we learned from not talking about data

17/09/2020 Revising the Caldicott Principles: our response

08/10/2020 We've grown

30/10/2020 Lessons from setting up the INSIGHT Data Trust Advisory Board

02/11/2020 Help us learn about health care professionals' attitudes to the use of health data

12/11/2020 The role of routine data in the fight against Covid-19

17/11/2020 Data partnerships and the pandemic: reflections from our parliamentary webinar

09/12/2020 What we mean by trustworthy use of patient data

17/12/2020 Our response to the National Data Strategy consultation

07/01/2021 Hello 2021

03/02/2021 How are stories about health data covered in the UK media?  
15/02/2021 Dame Fiona Caldicott  
17/02/2021 Debating data: a mini citizens' jury during a perfect storm  
23/02/2021 Engaging with Black and South Asian people on equitable data collection  
16/03/2021 UPDATE: working with patients and the public to explain access to health data  
16/03/2021 Working with patients and the public to explain access to health data  
15/04/2021 What counts as a 'public benefit' for data use?  
06/05/2021 New research and resources  
27/05/2021 Our perspective on the new system for GP data  
08/06/2021 We're looking for an evaluator for a new project  
15/06/2021 Trustworthy use of GP data: what must happen now  
01/07/2021 Listening to Black and South Asian people's views on health data  
05/08/2021 The future of Understanding Patient Data  
12/08/2021 Our recommendations for the data strategy for health and care  
02/09/2021 What we know about communicating how health data is used  
08/09/2021 The what and why of trusted research environments  
29/09/2021 Explaining the role of patient data in health charities guidance  
26/10/2021 What's next for Understanding Patient Data  
01/11/2021 Work with us to develop resources on datasets and pseudonymisation  
10/01/2022 Hello 2022  
31/01/2022 Equitable data collection: can you deliver the next stage?  
02/03/2022 New research: primary care professionals' views on the use of health data  
26/04/2022 New research: insights from Black and South Asian people on patient data  
31/05/2022 Deepening our understanding of barriers to equitable health data collection for Black and South Asian people  
09/06/2022 New resources: explaining patient data in an accessible way  
13/12/2022 Closing gaps in patient data for Black and South Asian communities