natureoutlook

BIG DATA IN BIOMEDICINE

5 November 2015 / Vol 527 / Issue No 7576



Cover art: Tatiana Plakhova

Editorial

Herb Brody Michelle Grayson Eric Bender Nick Haines Jenny Rooke

Art & Design

Wesley Fernandes Denis Mallet Annthea Lewis

Production

Karl Smart Ian Pope Mira Loufti

Sponsorship

George Sun Samantha Morley

Marketing

Hannah Phipps

Project Manager Anastasia Panoutsou

Art Director Kelly Buckheit Krause

Publisher

Richard Hughes

Magazine Editor Rosie Mestel

Editor-in-Chief

Philip Campbell

't may now cost less to sequence the three billion DNA base pairs of a human genome than to do a brain scan. But how does all that genomic data translate into treatment?

Life scientists are bringing together astonishing volumes of information from genomic sequencing, lab studies and patient records. And the resulting era of 'precision medicine' is already delivering treatments tailored to individual needs.

These 'big data' efforts face huge challenges, from creating analytic tools and solving scientific puzzles to accessing millions of gigabytes of data and overcoming barriers to accessing patients' health records (see pages S2 and S19).

Dozens of international projects are producing huge amounts of biomedical information, not just on the genome but on many other '-omes' (S8). Giant strides are being made in mapping the human proteome and building a 'parts list' of the body (S6). Meanwhile, smartphones and other wearable devices are generating continuous flows of health data from large numbers of people (S12). This vast array of data will allow a more detailed understanding of disease traits in analyses known as deep phenotyping (S14). Research organizations are assembling cloud-based 'information commons' to standardize, store and share the data (S16).

Drug companies are facing complex choices (S18). Many are opting to treat cancer, a main thrust in national programmes such as the UK 100,000 Genomes Project (S5). And some of these therapies are already changing clinical practice (S10).

We are pleased to acknowledge that this Outlook was produced with support from the National Center for Protein Sciences-Beijing, Beijing Proteome Research Center, State Key Laboratory of Proteomics, China Human Proteome Organization, Beijing Institute of Radiation, and the Academy of Military Medical Sciences. As always, Nature retains sole responsibility for all editorial content.

Eric Bender

Contributing Editor

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CITING THE OUTLOOK

Cite as a supplement to Nature, for example, Nature Vol. XXX, No. XXXX Suppl., Sxx-Sxx (2015).

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