

## Market Round-up

V22 – 19<sup>th</sup> October 2018

### General Market News:

- **Ger Brennan** and **MSD Ireland** have announced they are to create 170 new jobs in Carlow.  
[LINK](#)
- **HSE** secures lifting of contract suspension for the supply of **GSK's Infanrix Hex** of 6 in 1 vaccines.  
[LINK](#)
- **GSK** has appointed **Publicis Media** as its global media buying and planning agency. The account is estimated to be worth \$1.7bn.  
[LINK](#)
- **Shire** announces UCD master's scholarship in biopharmaceutical engineering  
[LINK](#)
- The **MMP** is currently undertaking a period of consultation in relation to their Roadmap for the prescribing of best-value biological (BVB) medicines (including biosimilars). Details of this consultation, including the draft document, can be found in the consultation section of the MMP website.  
[LINK](#)
- **Roche's Kadcyla** shows promise in early breast cancer during Phase III trials.  
[LINK](#)
- **AstraZeneca** and **MSD's** PARP inhibitor **Lynparza** has been granted orphan drug designation in the US for the treatment of pancreatic cancer.  
[LINK](#)
- **HSE** told to reinstate consultants' pay scales  
[LINK](#)
- **Pfizer** is cutting about 2% of its workforce by early 2019  
[LINK](#)
- **NICE** rejects **Eli Lilly's** breast cancer drug **Verzenio**  
[LINK](#)

### NCPE Drug Updates

- A full HTA is recommended to assess the clinical effectiveness of **Novartis's Kymriah**  
[LINK](#)
- A full HTA is recommended to assess the clinical effectiveness and cost effectiveness of **Gilead's Yescarta**  
[LINK](#)
- A full HTA is recommended to assess the clinical effectiveness and cost effectiveness of **AbbVie's Venclyxto**  
[LINK](#)

## Biologics / Biosimilars:

- "The **HSE** may miss out on €37m savings as costly drug **Humira** comes off patent as the policy of switching to biosimilar drugs was never adopted" *Irish Times article*  
[LINK](#)
- **AbbVie** settles **Humira** biosimilar patent disputes with **Novartis**  
[LINK](#)
- **Amgen** has begun launching **Amgevita**, the first biosimilar of **AbbVie's** anti-inflammatory blockbuster **Humira** to win approval in Europe, across the region.  
[LINK](#)
- **Samsung Bioepis** has announced the European launch of **Humira** biosimilar **Imraldi**, as NHS chiefs tell Trusts to ensure they are ready to realise the potential savings from using cheaper versions of the world's biggest selling drug.  
[LINK](#)
- **MDS** drops **Lantus** biosimilar, blames pricing and production cost concerns  
[LINK](#)

## Drug Approvals:

- **Pfizer** has received approval from the FDA for **Talzenna** (talazoparib), its PARP inhibitor for a genetically defined line of breast cancer.  
[LINK](#)

## Drug Approval Submissions:

- US and European regulators have accepted filings for **Novartis's siponimod** for secondary progressive multiple sclerosis (SPMS).  
[LINK](#)
- The FDA's Gastrointestinal Drugs Advisory Committee convened on Thursday, October 18, to review and discuss a marketing application from **Shire** for their constipation medicine **prucalopride**  
[LINK](#)
- **ViiV Healthcare** has submitted a New Drug Application to the FDA for their two-drug, single pill HIV treatment  
[LINK](#)

## Brexit:

- **Batch Testing** And Brexit: 4 Things You Need To Know  
[LINK](#)
- The UK Pharmaceutical Industry Braces for Brexit  
[LINK](#)
- Brexit: 'No optimism' for summit breakthrough, says Donald Tusk  
[LINK](#)

## Further Reading:

- Tufts study uncovers the economic advantage of single-source drug development and manufacturing  
[LINK](#)
  - Fighting The Opioid Epidemic: How Grünenthal's Abuse-Deterrent Technology Contributes  
[LINK](#)
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