

## Market Round-up

V14 – 24<sup>th</sup> August, 2018

### General Market News:

- **Uniphar** was cleared to buy **Sisk's** hospital equipment business for a for reported €65m  
[LINK](#)  
**HSE** under fire for hiring 600 extra senior managers  
[LINK](#)
- **Pfizer** has made an equity investment of an undisclosed size in immunotherapy firm **Biontech** through a multi-year research and development collaboration worth up to \$425m.  
[LINK](#)
- **NHS trusts** are warning that the health service is not prepared for a no-deal **Brexit** and that drug supplies could be affected as a result.  
[LINK](#)
- The early launch of **Amgen's** migraine drug **Aimovig** drug is likely to give it a solid head start over competitors.  
[LINK](#)
- **Amgen** to present new **Repatha** analyses at ESC Congress 2018  
[LINK](#)
- **AstraZeneca** inhaler lags **GSK** drug in lung disease trial  
[LINK](#)
- **Pfizer, Astellas** revise two cancer drug trial protocols to speed up results evaluating the safety and efficacy of **XTANDI® (enzalutamide)**  
[LINK](#) and [LINK](#)
- **Pfizer** loses bid to escape **Lipitor** antitrust lawsuit  
[LINK](#)

### Biosimilars:

- **Pfizer** lawsuit could clear path for biosimilars  
[LINK](#)

### NCPE Drug Updates

- No updates from NCPE this week.

### Drug Approvals:

- EU approves **Pfizer's Xeljanz** for **ulcerative colitis**  
[LINK](#)

- The **FDA** announced that it has given approval to **Teva Pharmaceuticals'** generic version of **EpiPen** and **EpiPen Jr.** (epinephrine) auto-injector. This is the first generic version of EpiPen and EpiPen Jr.  
[LINK](#)
- **Merck, Eisai** get **FDA** approval for **Lenvima** label expansion  
[LINK](#)
- **Merck** obtains earlier **FDA** approval for expanded label of **Keytruda** in front-line Lung cancer  
[LINK](#)
- **Shire's Takhzyro** has been approved in the US for the prevention of attacks of hereditary angioedema (**HAE**) in patients 12 years of age and older  
[LINK](#)

### **Brexit:**

- The UK has released its first set of guidelines on what to do if there's a no-deal Brexit. In 24 technical papers, the government outlines its contingency plans for industries including medicine in the event that negotiations should fail.  
[LINK](#)
- What to do if there is no Brexit deal. Article from *Financial Times*  
[LINK](#)
- The British government has told pharmaceutical companies to stockpile an extra six weeks supply of medicines to prepare for a no-deal Brexit and warned that businesses will face new bureaucratic hurdles when they import and export goods.  
[LINK](#)

### **Further Reading:**

- Researchers hail "exciting results" in ovarian, lung cancer trial  
[LINK](#)
- Direct-to-consumer genetic testing is booming, but ethical concerns remain  
[LINK](#)
- **Sláintecare** must not be sacrificed at the altar of Irish politics – article appeared in Irish Times.  
[LINK](#)
- Despite 70% of companies saying that Britain's exit from the EU will have a negative economic impact, just 6% of Irish businesses have a formal **Brexit** plan in place, according to a new report from **AIB**.  
[LINK](#)
- Machine learning is widely predicted to make drug discovery and patient diagnosis quicker, cheaper and more effective in the future, and signs of this can already be seen.  
[LINK](#)
- A U.S. appeals court on Wednesday struck out a \$3 million verdict against **GlaxoSmithKline** over the suicide of an attorney who took a **generic version** of the company's antidepressant **Paxil**, finding the company could not be held liable for injuries allegedly caused by a generic copy  
[LINK](#)