AstraZeneca
What science can do

Taking our science to the ASCO 2018 Annual Meeting

Investor and analyst planner and information pack

Chicago, Illinois, USA

01 June 2018

Forward-looking statements

In order, among other things, to utilise the 'safe harbour' provisions of the US Private Securities Litigation Reform Act 1995, we are providing the following cautionary statement:

This document contains certain forward-looking statements with respect to the operations, performance and financial condition of the Group, including, among other things, statements about expected revenues, margins, earnings per share or other financial or other measures. Although we believe our expectations are based on reasonable assumptions, any forward-looking statements, by their very nature, involve risks and uncertainties and may be influenced by factors that could cause actual outcomes and results to be materially different from those predicted. The forward-looking statements reflect knowledge and information available at the date of preparation of this document and AstraZeneca undertakes no obligation to update these forward-looking statements. We identify the forward-looking statements by using the words 'anticipates', 'believes', 'expects', 'intends' and similar expressions in such statements. Important factors that could cause actual results to differ materially from those contained in forward-looking statements, certain of which are beyond our control, include, among other things: the loss or expiration of, or limitations to, patents, marketing exclusivity or trademarks, or the risk of failure to obtain and enforce patent protection; effects of patent litigation in respect of IP rights; the impact of any delays in the manufacturing, distribution and sale of any of our products; the impact of any failure by third parties to supply materials or services; the risk of failure of outsourcing; the risks associated with manufacturing biologics; the risk that R&D will not yield new products that achieve commercial success; the risk of delay to new product launches; the risk that new products do not perform as we expect; the risk that strategic alliances and acquisitions, including licensing and collaborations, will be unsuccessful; the risks from pressures resulting from generic competition; the impact of competition, price controls and price reductions; the risks associated with developing our business in emerging markets; the risk of illegal trade in our products; the difficulties of obtaining and maintaining regulatory approvals for products; the risk that regulatory approval processes for biosimilars could have an adverse effect on future commercial prospects; the risk of failure to successfully implement planned cost reduction measures through productivity initiatives and restructuring programmes; the risk of failure of critical processes affecting business continuity; economic, regulatory and political pressures to limit or reduce the cost of our products; failure to achieve strategic priorities or to meet targets or expectations; the risk of substantial adverse litigation/government investigation claims and insufficient insurance coverage; the risk of substantial product liability claims; the risk of failure to adhere to applicable laws, rules and regulations; the risk of failure to adhere to applicable laws, rules and regulations relating to anti-competitive behaviour; the impact of increasing implementation and enforcement of more stringent anti-bribery and anti-corruption legislation; taxation risks; exchange rate fluctuations; the risk of an adverse impact of a sustained economic downturn; political and socio-economic conditions; the risk of environmental liabilities; the risk of occupational health and safety liabilities; the risk associated with pensions liabilities; the impact of failing to attract and retain key personnel and to successfully engage with our employees; the risk of misuse of social medial platforms and new technology; and the risk of failure of information technology and cybercrime. Nothing in this presentation / webcast should be construed as a profit forecast.

Key AstraZeneca abstracts

Date	Time	Abstract #	Title
Friday 1 June			
	14:45	2503	Molecular analysis for therapy choice (MATCH) arm W: Phase II study of AZD4547 in patients with tumors with aberrations in the FGFR pathway
	16:30	6011	Neoadjuvant anti-OX40 (MEDI6469) prior to surgery in head and neck squamous cell carcinoma
Saturday 2 June			
	15:00	7004	Moxetumomab pasudotox in heavily pretreated patients with relapsed/refractory hairy cell leukemia: Results of a pivotal international study
	!5:00	10503	SPRINT: Phase II study of the MEK 1/2 inhibitor selumetinib (AZD6244, ARRY-142886) in children with neurofibromatosis type 1 (NF1) and inoperable plexiform neurofibromas (PN)
Sunday 3 June			
	08:00	1001 / 1002	Genetic landscape of resistance to CDK4/6 inhibition in circulating tumor DNA (ctDNA) analysis of the PALOMA3 trial of palbociclib and fulvestrant (*Faslodex*) versus placebo and fulvestrant (*Faslodex*) / Abemaciclib for pre/perimenopausal women with HR+, HER2- advanced breast cancer
	08:00	1007	AZD5363 plus paclitaxel versus placebo plus paclitaxel as first-line therapy for metastatic triple-negative breast cancer (PAKT): A randomised, double-blind, placebo-controlled, phase II trial
	09:45	104	Randomised phase II neoadjuvant study (GeparNuevo) to investigate the addition of durvalumab (*Imfinzi*) to a taxane-anthracycline containing chemotherapy in triple negative breast cancer (TNBC)
	09:45	7501	Acalabrutinib (*Calquence)* in patients (pts) with Waldenström macroglobulinemia (WM)
Monday 4 June			
	08:00	8503	DREAM: A phase II study of durvalumab (*Imfinzi*) with first line chemotherapy in mesothelioma—First results
	08:00	502	Role of adding ovarian function suppression to tamoxifen (*Nolvadex*) in young women with hormone-sensitive breast cancer who remain premenopausal or resume menstruation after chemotherapy: The ASTRRA study
	15:00	5003	Study 08 Olaparib (*Lynparza*) combined with abiraterone in patients (pts) with metastatic castration-resistant prostate cancer (mCRPC): A randomized phase II trial
	15:00	LBA4008	Chemoprevention of esophageal cancer with esomeprazole (*Nexium*) and aspirin therapy: Efficacy and safety in the phase III randomized factorial ASPECT trial
Tuesday 5 June			
	09:45	5504	Phase I trial of olaparib (*Lynparza*, PARP inhibitor) and vistusertib (mTORC1/2 inhibitor) in recurrent endometrial, ovarian and triple negative breast cancer

Investor event

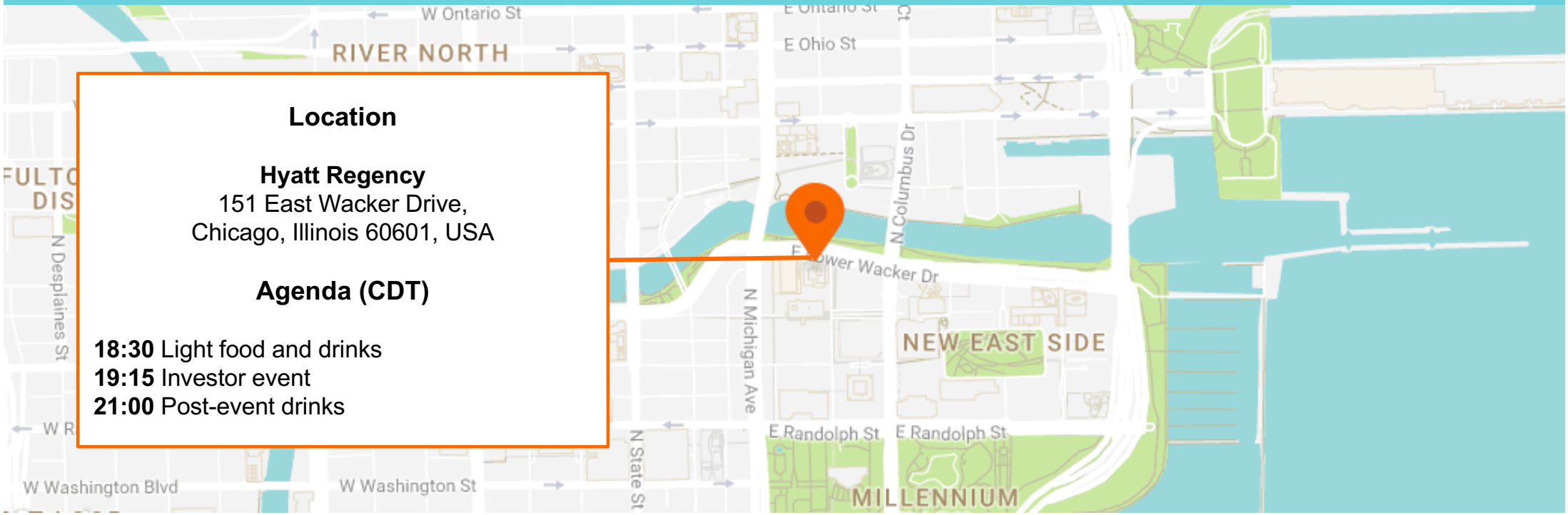
4 June 2018

Location

Hyatt Regency
151 East Wacker Drive,
Chicago, Illinois 60601, USA

Agenda (CDT)

18:30 Light food and drinks
19:15 Investor event
21:00 Post-event drinks



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