AGENDA OVERVIEW

	E	uropean Antik	oody Congress			-World-Immunot	herapy Congress		World Biosimi	ilar Congress	_Clinical Trial Wo	rkshop Day 1 & 2
		iaropouri Arraik	Jour Congress				29th October 2018		World Blocking	ilai Sengiese	Cililical IIIal III	Thomps buy i a 2
Morning plenary keynote						Morning plenary keynote			Morning plen	nary keynote	Targeted workshops focusing on how we can reduce trial	
					Morning netv	vorking break					failure and improve patient e Transforming clinical trials to	meet demands of the futur
Plenary roundtables						Plenary roundtables			Plenary ro	undtables	Case studies on disruptive technology within clinical in the clinical in	
					Lur	nch						
Protein engineering and R&D Engineering antibody formats Display of biologics Bispecifics development • Investor panel • Regulatory updates • Antibody naming		nel updates	discoveryADC and loadedIrF		vestment and regulation Investor panel Regulatory updates Antibody naming	Immune checkpoint inhibitors • Approved Market Formats • New Agents • Combination Agents	Antigen Targets Cell Therapy for solid tumours Allogeneic gene therapy The target served to the		Market access strategies, opportunities and commercial challenges • Ensuring clinical adoption • Bringing biosimilars to market • Technology transfer • Emerging markets	Biosimilar Development Manufacture and clinical case studies Clinical trials Demonstrating biosimilarity Opportunities in development		
Closing keynote						Closing keynote			Closing keynote		HPAPI World Congress	
						Day 2 – Tuesday 3	30th October 2018					
Opening keynotes: C-level panel					Opening keynotes: C-level panel			Opening keynotes: C-level panel		Opening keynotes		
						Morning re	efreshments					
Protein engineering and R&D • Antibody screening and analytics	development / discovery		Armed antibodies discovery / development Novel payloads Al in discovery / development Articles And engineering		Platform technology showcase • New technology for antibody discovery and development	Immune checkpoint inhibitors • Approved Market Formats • New Agents • Combination Agents	 Cell therapy Antigen Targets Cell Therapy for solid tumours Allogeneic gene therapy 	opportunities and commercial challenges Regulatory Updates Streamlining approval		Biosimilar Development Demonstrating interchangeability Analytics, development & pharmacovigilance Harmonizing immunogenicity testing Characterization, assay development & extrapolation	Roundtables/workshops Considerations for risk assessments EHS challenges of new ADCs Toxicological issues Outsourcing Adapting to guidelines Containment systems: old vs new	
						Lui	nch					
MC and evelopability Antibody commercialisation and manufacture Protein production Bispecifics discove expection of the support of the suppo		development • ADCs and armed antibodies in analytic preclinical and discovery • Al and i analytic new use		ery	Platform technology showcase • Antibody characterisation • Antibody manufacture	Market access Investor input Regulatory issues Commercialisation and manufacture	Clinical trials Workshops Challenges with clinical trials Case studies	opp con	arket access strategies, portunities and mmercial challenges P and legal considerations	Biosimilar Development Development & Manufacturing strategies	Containment, equipment, facility design Containment technologies Adapting solutions for HPAPIS	Occupational EH&S Toxicological limits Control banding Using data
Closing keynotes					Closing keynotes			Closing keynotes		Closing keynotes		
						Day 3 – Wednesday	y 31st October 2018					
Novel targets and biomarker discovery Early stage biomarker discovery Target development New biomarker development		purification technologies developm • New screening and analytical • Antibody		Updates or development	ngineering and optimising	Biomarkers Technology Development New biomarkers in development Immunotherapy for solid tumours Challenges for solid tumours Examples and pre/clinical development	and cancer vaccinesEarly stage discoveryTechnological				Outsourcing and manufacturing • Evolving vendor relationships • Supply chain integration • Case studies • Manufacturing processes Drug safety and pharmacovigilance • Current development • Challenges	Guidelines and regulations Aseptic facility design Cleaning requirement Update: EMA guidelin Antibody drug conjug Site specific conjugat Case studies
						Lu	nch					
		Closing	plenary			Closing	plenary				Closing	nlenary











