

香港中文大學醫學院 Facultý Of Medicine

The Chinese University Of Hong Kong



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Joint Chinese University of Hong Kong-New Territories East Cluster Clinical Research Ethics Committee

香港中文大學-新界東醫院聯網 臨床研究倫理 聯席委員會

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8/F, Lui Che Woo C Tel: (852) 3505 3935 / 2144 59	linical Sciences Building, Prince of 26 Fax: (852) 2646 6653	of Wales Hospital, Shatin, HK Website : http://www.crec.cuhk.edu.hk
The Joint CUHK-NTEC CREC is an and scientific review and oversight of operating procedure and the principle	clinical studies within the jurisdiction	CUHK/NTEC and authorized to perform ethics of CUHK/NTEC in accordance with its standard CH Good Clinical Practice.
CREC Ref. No.: 2015.723		13 JUN 'I'
To: Prof. Kai Shun LEUN Dept. of Ophthalmolo Visual Sciences Hong Kong Eye Hosp	ogy and	
This notice is issued by the Jo you, being the principal invest	int CUHK-NTEC CREC with gator of the following study at	respect to the application/submission by your study site:
• Study Protocol Title:	to guide intraocular pressur hypertensives (OHT)	r layer (RNFL) thinning as a biomarker e (IOP) lowering treatment in ocular
• Investigator(s):	Kai Shun LEUNG, Poemen P	ui Man CHEN and Felix Chi Hong LI
In accordance with our standareview of your application/sub	ard operating procedure, we hamission as detailed below:	ave duly performed ethics and scientific
 Nature of Your Application/Submission: 	☐ Initial application ☐ Amendments/changes	☐ Others: ☐ Renewal
• Mode of Review:	☐ Full review	Expedited review
 Date of Initial/Renewal Approval: 	09 March 2017	
 Date of Amendment Approval: 	01 June 2017	
• Document(s) Reviewed:	See Schedule 1	
• Reviewer(s):	See Schedule 2	
After due review by our re application/submission as fol	viewer(s), we hereby write to lows:	o inform you of our decision on your
• Decision:	condition(s) below) Application/Submission below)	approved approved with condition(s) (see approved with remark(s) (see remark(s) approved with condition(s) and

• Condition(s):

A copy of the Certificate for Clinical Trial is required to be submitted to the Joint CUHK-NTEC CREC prior to commencement of the study

remark(s) (see condition(s) and remark(s) below)



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Tel: (852) 3505 3935 / 2144 5926

Fax: (852) 2646 6653

Website: http://www.crec.cuhk.edu.hk

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• Regular Progress Report(s) Required: Every 12 months from the date of initial/renewal approval and during the period of the study if required

You, being the principal investigator of the study at your study site, are reminded to comply with our requirements and to maintain communication with us during the period of the study by undertaking the principal investigator's responsibilities including (but not limited to):

- if the study is an industry-sponsored clinical study, submitting to us a copy of the fully executed indemnity agreement satisfying the Hospital Authority's requirement prior to commencement of the study (if it has not been submitted yet);
- observing and complying with all applicable requirements under our standard operating procedure ("IRB/REC SOP"), the Declaration of Helsinki and the ICH GCP (if applicable);
- submitting regular progress report(s) at the required intervals (as specified above) in accordance with the requirements in the IRB/REC SOP;
- not implementing any amendment/change to any approved study document/material without our written approval, except where necessary to eliminate any immediate hazard to the subjects or if an amendment/change is only of an administrative or logistical nature;
- notifying us of any new information that may adversely affect the rights, safety or well-being of the subjects or the proper conduct of the study;
- reporting any deviation from the study protocol or compliance incident that has occurred during the study and may adversely affect the rights, safety or well-being of any subject in accordance with the requirements in the IRB/REC SOP;
- submitting safety reports on all SAEs observed at your study site or SUSARs reported from outside your study site in accordance with the requirements in the IRB/REC SOP; and
- submitting a final report in accordance with the requirements in the IRB/REC SOP upon completion or termination of the study at your study site.

In addition to the above, you are also reminded to observe and comply with other applicable regulatory and management requirements including (but not limited to):

- if required by Hong Kong laws or regulations, obtaining a certificate for clinical trial through the Hong Kong Department of Health and complying with the associated requirements;
- obtaining the necessary consent from the management of your institution/department in accordance with the requirements of your institution/department;
- if required by local laws or regulations at conducting site out of IRB/REC's jurisdiction, obtaining an approval and complying with associated requirements;
- not representing to any third party or in any way likely to mislead any third party forming the view that the approval from the IRB/REC has any extraterritorial effect; and
- with due diligence ensuring your teams, staff, agents or whosoever connected with you to comply with the preceding requirements.

Yours sincerely,

Envy Lee (Secretary)

for and on behalf of

The Joint CUHK-NTEC CREC

Schedule 1 Documents Reviewed

The documents reviewed by with respect to the said application/submission include:

Amendment dated 24 January 2017

- Research Protocol, Version 2, dated 24 January 2017 (clean and track changes version)
- Participant Information Sheet and Informed Consent Form, English Version 2, dated 24 January 2017 (clean and track changes version)
- Participant Information Sheet and Informed Consent Form, Chinese Version 2, dated 24 January 2017 (clean and track changes version)

Schedule 2 Reviewers List Joint CUHK-NTEC Clinical Research Ethics Committee

Title and Name	Occupation	Qualification	Male / Female (M/F)
Prof. Simon LAM	Associate Professor, Department of Paediatrics, CUHK	MBBChir, DCH(HK), DCH(International), MRCPCH, FHKCPaed, FHKAM(Paed)	М
Dr. Keary ZHOU	Instructor, School of Pharmacy, CUHK	BS(UCLA), PharmD(USC)	F





Joint Chinese University of Hong Kong-New Territories East Cluster Clinical Research Ethics Committee

香港中文大學-新界東醫院聯網 臨床研究倫理 聯席委員會

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Tel: (852) 2632 3935 / 2144 5926 Fax: (852) 2646 6653 Website: http://www.crec.cuhk.edu.hk

The Joint CUHK-NTEC CREC is an independent committee established by CUHK/NTEC and authorized to perform ethics and scientific review and oversight of clinical studies within the jurisdiction of CUHK/NTEC in accordance with its standard operating procedure and the principles of the Declaration of Helsinki and ICH Good Clinical Practice.

	ciples of the Declaration of Helsinki and ICH Good Clinical Practice.	
CREC Ref. No.: 2015.72	23 Z2 MGR	516
To: Prof. Kai Shun LE Dept. of Ophthalm Visual Sciences Hong Kong Eye H	nology and	
	e Joint CUHK-NTEC CREC with respect to the application/submission by estigator of the following study at your study site:	
• Study Protocol Title:	Progressive retinal nerve fiber layer (RNFL) thinning as a biomarker to guide intraocular pressure (IOP) lowering treatment in ocular hypertensives (OHT)	
• Investigator(s):	Kai Shun LEUNG, Poemen Pui Man CHEN and Felix Chi Hong LI	
	ndard operating procedure, we have duly performed ethics and scientific submission as detailed below:	
• Nature of Your Application/Submission	☑ Initial application☐ Others:☐ Amendments/changes☐ Renewal	
• Mode of Review:		
• Date of Initial/Renewal Approval:	09 March 2016	
• Document(s) Reviewed:	See Schedule 1	
• Reviewer(s):	See Schedule 2	
After due review by our application /submission as f	reviewer(s), we hereby write to inform you of our decision on your follows:	
• Decision:	 □ Application/Submission approved ⋈ Application/Submission approved with condition(s) (see condition(s) below) □ Application/Submission approved with remark(s) (see remark(s) below) □ Application/Submission approved with condition(s) and remark(s) (see condition(s) and remark(s) below) 	
• Condition(s):	A copy of the Certificate for Clinical Trial is required to be submitted to the Joint CUHK-NTEC CREC prior to commencement of the study	
• Regular Progress Report(s) Required:	Every 12 months from the date of initial/renewal approval and during the period of the study if required	
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- submitting regular progress report(s) at the required intervals (as specified above) in accordance with the requirements in the IRB/REC SOP;
- not implementing any amendment/change to any approved study document/material without our written approval, except where necessary to eliminate any immediate hazard to the subjects or if an amendment/change is only of an administrative or logistical nature;
- notifying us of any new information that may adversely affect the rights, safety or well-being of the subjects or the proper conduct of the study;
- reporting any deviation from the study protocol or compliance incident that has occurred during the study and may
 adversely affect the rights, safety or well-being of any subject in accordance with the requirements in the IRB/REC
 SOP.
- submitting safety reports on all SAEs observed at your study site or SUSARs reported from outside your study site in accordance with the requirements in the IRB/REC SOP; and
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- obtaining the necessary consent from the management of your institution/department in accordance with the requirements of your institution/department;
- if required by local laws or regulations at conducting site out of IRB/REC's jurisdiction, obtaining an approval and complying with associated requirements;
- not representing to any third party or in any way likely to mislead any third party forming the view that the approval from the IRB/REC has any extraterritorial effect; and
- with due diligence ensuring your teams, staff, agents or whosoever connected with you to comply with the preceding requirements.

Yours sincerely,

Jenny Ng (Secretary) for and on behalf of

The Joint CUHK-NTEC CREC

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Schedule 1 Documents Reviewed

The documents reviewed by with respect to the said application/submission include:

- Research Protocol, Version 1, dated 18 January 2016
- Participant Information Sheet and Informed Consent Form, English Version 1, dated 18 January 2016
- Participant Information Sheet and Informed Consent Form, Chinese Version 1, dated 18 January 2016

Schedule 2 Reviewers List Joint CUHK-NTEC Clinical Research Ethics Committee

Title and Name	Occupation	Qualification	Male / Female (M/F)	Study Reviewed by	Present in CREC meeting on 05 Jan 2016
Chairman: Prof. Benny C.Y. ZEE	Professor, School of Public Health, CUHK	BSc(Manitoba), MSc(Manitoba), PhD(Pittsburgh)	M	~	٧
Vice/Deputy Chairman: Dr. Chi Kong LI	Consultant Paediatrician, Department of Paediatrics, PWH	MBBS, MD(CUHK), MRCP(UK), DCH(Lond), FHKCPaed, FHKAM(Paed), FRCPCH(UK)	M	. 1	٨
Prof. Lijia CHEN	Assistant Professor, Department of Ophthalmology Visual Sciences, CUHK	Bachelor of Med (China), Master of Op (China), PhD (CUHK), FRCS (Edin), MRCS (Edin)	М	4	
Prof. Alice Pik Shan KONG	Associate Professor, Department of Medicine and Therapeutics, CUHK	MBChB(CUHK), MRCP(UK), FHKCP, FHKAM(Medicine), FRCP(Glasgow)	F		
Prof. Bonnie Ching Ha KWAN	Associate Professor, Department of Medicine and Therapeutics, PWH, CUHK	MBBS(London), MHKCP, MRCP(UK), FHKCP, FHKAM	F		
Prof. Brigette MA	Professor, Department of Clinical Oncology, CUHK	FRACP(Australia),F HKCP, FHKAM(Medical Oncology)	F		
Prof. Vincent C.T. MOK	Associate Professor, Department of Medicine and Therapeutics, CUHK	MBBS(U Sydney), MRCP(UK), FHKCP, FHKAM, MD(CUHK), FRCP(Edin)	M		
Prof. Cheuk Chun SZETO	Professor, Department of Medicine and Therapeutics, CUHK	MBChB(CUHK), MRCP(UK), FHKCP, FHKAM, DM(CUHK), FRCP(Edin)	М	4	4
Prof. Wai Kwong TANG	Professor, Department of Psychiatry, CUHK	MBChB(CUHK), MD (CUHK), MRCP(UK), FHKCP, FHKAM	M	1	4
Prof. Brian TOMLINSON	Professor, Department of Medicine and Therapeutics, CUHK	BSc, MBBS, MD, MRCP(UK), FHKCP, FRCP, FRC(E), FRCP(G), FHKAM(Med), FCP, FACP	· M		1

Title and Name	Occupation	Qualification	Male / Female (M/F)	Study Reviewed by	Present in CREC meeting on 05 Jan 2016
Dr. Simon K.C. CHAN	Consultant, Department of Anaesthesia and Intensive Care, PWH	MBBS (UNSW), FANZCA(Aust), FHKCA, FHKAM, Dip Pain Mgt(HKCA), MHSM(UNSW)	M	√	√
Dr. Ernest H.M. MA	Senior Medical Officer, Department of Medicine and Geriatrics, TPH	MBChB, MRCP(UK), Msc Resp Med(Lond), FRCP(Lond, Edin, Ire), FHKCP, FHKAM, MBA, EdD(UTS)	М		
Dr. Kevin Ka Hang OR	Consultant, Department of Medicine and Geriatrics, SH	MBBS(UNSW), MRCP(UK), FHKCP, FHKAM(Med), FRCP(Edin), BChinMed(HKU)	М		
Dr. Keary ZHOU	Instructor, School of Pharmacy, CUHK	BS(UCLA), PharmD(USC)	F	٧	
Mr. Bryan Ping Ho CHUNG	Physiotherapist I, Physiotherapy Department, TPH	BSc in Physiotherapy, MSc in Health Care (Physiotherapy)	М	1	
Ms. Alexandra Dak Wai LO	Chinese Medicine Practitioner, Part-time lecturer and adjunct tutor, Dept of Anatomical and Cellular Pathology, PWH, CUHK	LLB, Hons.(HKU), PCLL(HKU), LLB, Hons.(Peking), LLM (CityU), BchinMed(HKU)	F		
Ms. Emily May Ling CHAN	Retired	DSW(HKPU), RSW Certified Hypnotherapist CISM, UMBC	F		٧
Mr. Christopher LIU	Executive Director, Liu Chong Hing Investment Ltd.	Bachelor of Arts (Oxford), Master of Arts in Jurisprudence (Oxford)	M	V	V
Mr. Wilson SO	Retired	Bachelor of Social Science, Master of Town and Country Planning	М	1	√
Mr. Ping Hei TAO	Retired	MHRM(MQU), Dip Soc Sci(HKBU)	М		
Mr. Foster YIM	Barrister-at-Law	PCLL(CUHK), JD(CUHK), Msc in Marketing (CUHK),MA in Philosophy (UK), BA (Hons) in Translation (LU)	М		