



Robert Mijumbi,
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13th August 2021

The Executive Secretary,
 National Drug Authority,
 Rume Towers, Plot 19 Lumumba Avenue,
 P.O. Box 23096, Kampala.

Dear Scientist,
REQUEST FOR PERMISSION TO CONDUCT PHASE I CLINICAL TRIAL OF MY COVID-19 CURE CANDIDATE (BertoCOV)

As you must have known through my previous letters to you, I have been conducting investigations to develop therapeutics for HIV, Hepatitis B, Cancer and Covid-19. Basing on my laboratory results, the Covid-19 cure candidate is ripe for phase I clinical trial and I request that the National Drug Authority give me clearance to do so, starting with the oral regimen. Attached to this letter is a profile of Covid-19 cure candidate, and for ease of future reference, those of HIV, Hepatitis B and Cancer are also attached. Also, a one (1) gram sample of each of the candidates is attached and marked as follows; HIV (A), Hepatitis B (B), Cancer (C), and Covid-19 (D).

Sincerely,

[Signature]
 Robert Mijumbi
 C.E.O. Bbert Research Group Ltd.



- CC: H.E. Gen. Yoweri Kaguta Tibuhaburwa Museveni, President of the Republic of Uganda
 The Rt. Hon. Speaker, Parliament of the Republic of Uganda
 The Rt. Hon. Prime Minister
 The Director Research, Ministry of Science, Technology and Innovation
 The Permanent Secretary, Ministry of Health
 The Executive Secretary, Uganda National Council for Science and Technology
 The Pharmaceutical Society of Uganda
 The Uganda Medical Association
 The Uganda Medical and Dental Practitioners Council
 The Uganda Cancer Institute
 The Uganda Aids Commission
 The Inspector General of Police

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Amijumbi
13/08/2021



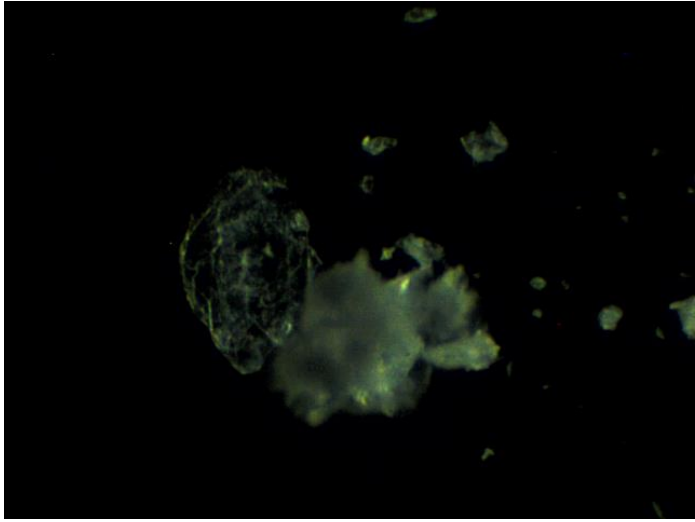
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Dept.	SA



PROFILE OF COVID-19 CURE CANDIDATE

Physicochemical Characteristics		
Dark Field Micrograph	Brand Name	BertoCOV
 <p style="text-align: right; margin-right: 50px;">X100</p>	Nature of API	Polypeptide
	Minimum Anticipated Biological Effect Level	660 mg
	Infrared Spectroscopy	See pages 9 & 10
	Composition of Processed Drug (%)	<ul style="list-style-type: none"> API 13.44% Excipients 86.56%
	Kinase Activity	None
	Stability	Biological function lost beyond 43°C
	Dissolution	100%
	Bioavailability	100%
	Dosage Form	Powder
	Route of Administration	Oral

Pharmacokinetics

The whole powder is dissolved in potable water at room temperature and drunk all at once. It is absorbed through the villi in the gastrointestinal tract and presented into the blood stream. It is progressively metabolized in the liver and then excreted renally.

Pharmacodynamics

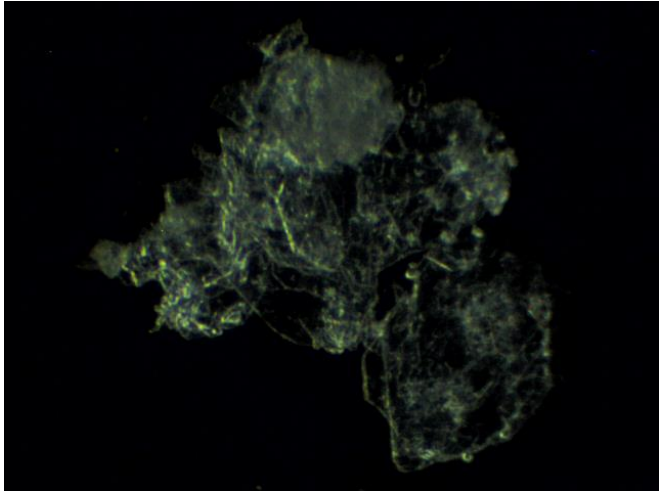
Once in the blood, it is circulated to various tissues and organs. It never penetrates individual cells but remains within the circulatory system where it digests the envelope of a viral particle before the virus infects a new cell. This exposes the viral enzymes necessary for replication to an unusual environment (plasma) and thus they cannot function; moreover, exposing the viral genome to plasma leads to its metabolism in the usual biochemical fashion.

In Vitro Interaction of API with Some of the Common Drugs

Category	Generic/ Trade Name	Effect on API Potency	
		Inhibitory	None
Antibiotic/ Antimicrobial	Amoxicillin		✓
	Ampicillin		✓
	Ampiclox (Ampicillin and Cloxacillin)		✓
	Ciprofloxacin	✓	
	Metronidazole		✓
	Erythromycin		✓
	Fluconazole		✓
	Septin		✓
Antimalarial/ Antiparasitic	Acyclovir		✓
	Fansida		✓

	Quinine	✓	
	Artemether- Lumefantrine		✓
	Albendazole		✓
Analgesic	Paracetamol		✓
	Paracetamol Caffeine Aspirin	✓	
	Diclofenac		✓
	Ibuprofen		✓
	Tramadol		✓
	Piroxicam		✓
Antihypertensive	Nifedipine		✓
	Losartan		✓
	Atenolol		✓
	Furosemide (Lasix)		✓
	Propranolol	✓	
Antidiabetic	Metformin		✓
	Glibenclamide		✓
Antiasthmatic	Aminophylline		✓
	Salbutamol		✓
	Prednisolone		✓
Anti-allergic	Cetirizine	✓	
	Cimetidine		✓
Antihyperlipidaemic	Atorvastatin		✓
Anticoagulant	Cardiac Aspirin		✓
Minerals	Magnesium/ Aluminium		✓
	Zinc	✓	
	Vitamin C		✓
	Vitamin B ₁ , B ₂ , B ₆ and Niacinamide		✓

PROFILE OF HIV CURE CANDIDATE

Physicochemical Characteristics		
Dark Field Micrograph	Brand Name	BertoV ₁
 <p style="text-align: right;">X100</p>	Nature of API	Polypeptide
	Minimum Anticipated Biological Effect Level	460 mg
	Infrared Spectroscopy	See pages 9 & 10
	Composition of Processed Drug (%)	<ul style="list-style-type: none"> • API 11.33% • Excipients 88.67%
	Kinase Activity	None
	Stability	Biological function lost beyond 43°C
	Dissolution	100%
	Bioavailability	100%
	Dosage Form	Powder
	Route of Administration	Oral

Pharmacokinetics

The whole powder is dissolved in potable water at room temperature and drunk all at once. It is absorbed through the villi in the gastrointestinal tract and presented into the blood stream. It is progressively metabolized in the liver and then excreted renally.

Pharmacodynamics

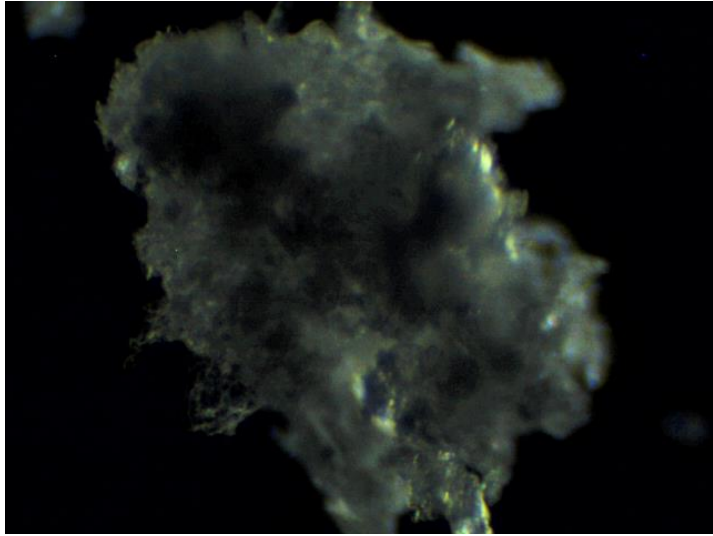
Once in the blood, it is circulated to various tissues and organs. It never penetrates individual cells but remains within the circulatory system where it digests the envelope of a viral particle before the virus infects a new cell. This exposes the viral enzymes necessary for replication to an unusual environment (plasma) and thus they cannot function. Also, it digests the viral genome; moreover, exposing the viral genome to plasma leads to its metabolism in the usual biochemical fashion.

In Vitro Interaction of API with Some of the Common Drugs

Category	Generic/ Trade Name	Effect on API Potency	
		Inhibitory	None
Antibiotic/ Antimicrobial	Amoxicillin		✓
	Ampicillin		✓
	Ampiclox (Ampicillin and Cloxacillin)		✓
	Ciprofloxacin	✓	
	Metronidazole		✓
	Erythromycin		✓
	Fluconazole		✓
	Septrin		✓
Antimalarial/ Antiparasitic	Acyclovir		✓
	Fansida		✓

	Quinine	✓	
	Artemether- Lumefantrine		✓
	Albendazole		✓
Analgesic	Paracetamol		✓
	Paracetamol Caffeine Aspirin	✓	
	Diclofenac		✓
	Ibuprofen		✓
	Tramadol		✓
	Piroxicam		✓
Antihypertensive	Nifedipine		✓
	Losartan		✓
	Atenolol		✓
	Furosemide (Lasix)		✓
	Propranolol	✓	
Antidiabetic	Metformin		✓
	Glibenclamide		✓
Antiasthmatic	Aminophylline		✓
	Salbutamol		✓
	Prednisolone		✓
Anti-allergic	Cetirizine	✓	
	Cimetidine		✓
Antihyperlipidaemic	Atorvastatin		✓
Anticoagulant	Cardiac Aspirin		✓
Minerals	Magnesium/ Aluminium		✓
	Zinc	✓	
	Vitamin C		✓
	Vitamin B ₁ , B ₂ , B ₆ and Niacinamide		✓

PROFILE OF HEPATITIS B CURE CANDIDATE

Physicochemical Characteristics		
Dark Field Micrograph	Brand Name	BertoHEP
 <p style="text-align: right; margin-right: 50px;">X100</p>	Nature of API	Polypeptide
	Minimum Anticipated Biological Effect Level	460 mg
	Infrared Spectroscopy	See pages 9 & 10
	Composition of Processed Drug (%)	<ul style="list-style-type: none"> • API 11.33% • Excipients 88.67%
	Kinase Activity	None
	Stability	Biological function lost beyond 43°C
	Dissolution	100%
	Bioavailability	100%
	Dosage Form	Powder
	Route of Administration	Oral

Pharmacokinetics

The whole powder is dissolved in potable water at room temperature and drunk all at once. It is absorbed through the villi in the gastrointestinal tract and presented into the blood stream. It is progressively metabolized in the liver and then excreted renally.

Pharmacodynamics

Once in the blood, it is circulated to various tissues and organs. It never penetrates individual cells but remains within the circulatory system where it digests the envelope of a viral particle before the virus infects a new cell. This exposes the viral enzymes necessary for replication to an unusual environment (plasma) and thus they cannot function. Also, it digests the viral genome; moreover, exposing the viral genome to plasma leads to its metabolism in the usual biochemical fashion.

In Vitro Interaction of API with Some of the Common Drugs

Category	Generic/ Trade Name	Effect on API Potency	
		Inhibitory	None
Antibiotic/ Antimicrobial	Amoxicillin		✓
	Ampicillin		✓
	Ampiclox (Ampicillin and Cloxacillin)		✓
	Ciprofloxacin	✓	
	Metronidazole		✓
	Erythromycin		✓
	Fluconazole		✓
	Septtrin		✓
Antimalarial/ Antiparasitic	Acyclovir		✓
	Fansida		✓

	Quinine	✓	
	Artemether- Lumefantrine		✓
	Albendazole		✓
Analgesic	Paracetamol		✓
	Paracetamol Caffeine Aspirin	✓	
	Diclofenac		✓
	Ibuprofen		✓
	Tramadol		✓
	Piroxicam		✓
Antihypertensive	Nifedipine		✓
	Losartan		✓
	Atenolol		✓
	Furosemide (Lasix)		✓
	Propranolol	✓	
Antidiabetic	Metformin		✓
	Glibenclamide		✓
Antiasthmatic	Aminophylline		✓
	Salbutamol		✓
	Prednisolone		✓
Anti-allergic	Cetirizine	✓	
	Cimetidine		✓
Antihyperlipidaemic	Atorvastatin		✓
Anticoagulant	Cardiac Aspirin		✓
Minerals	Magnesium/ Aluminium		✓
	Zinc	✓	
	Vitamin C		✓
	Vitamin B ₁ , B ₂ , B ₆ and Niacinamide		✓

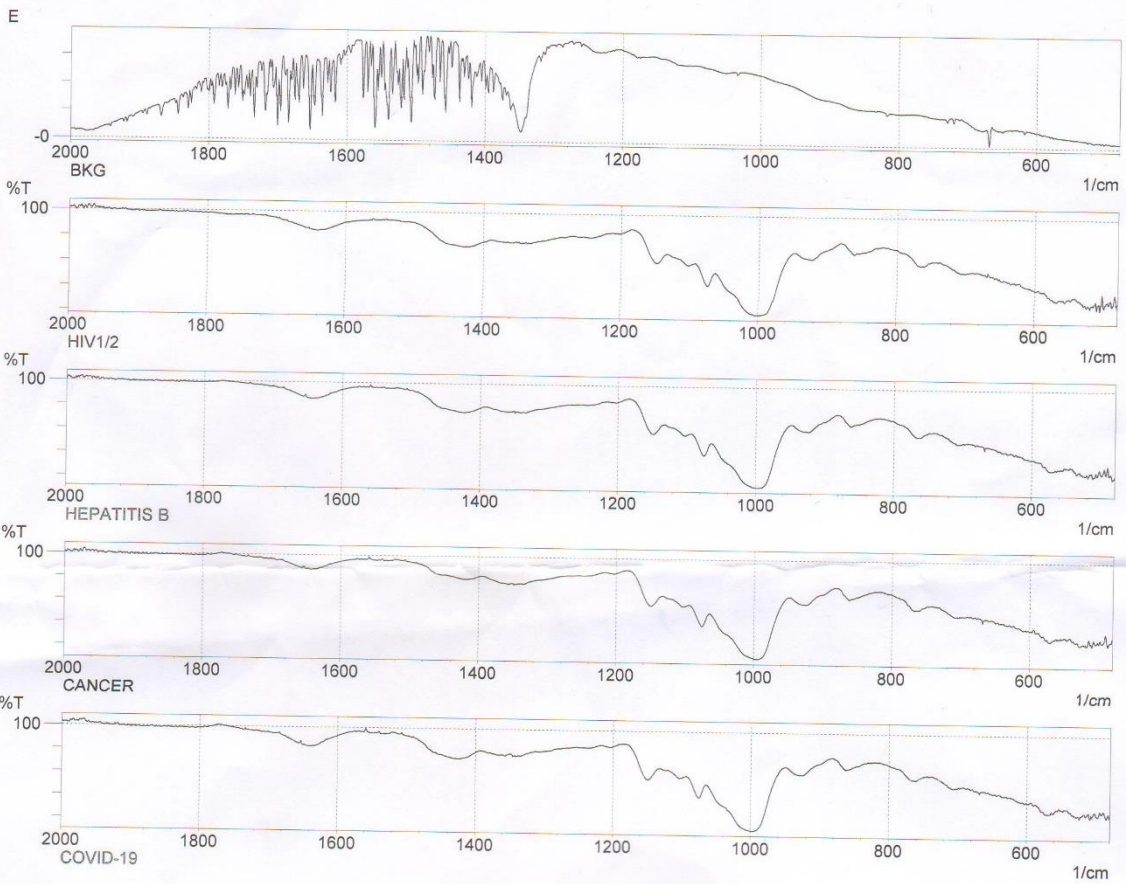
PROFILE OF CANCER CURE CANDIDATE

Physicochemical Characteristics			
Brand Name	BertoCAN		
Nature of API	Polypeptide		
Minimum Anticipated Biological Effect Level	110 mg		
Infrared Spectroscopy	See pages 9 & 10		
Composition of Processed Drug (%)	<ul style="list-style-type: none"> • API 5.16% • Excipients 94.84% 		
Kinase Activity	None		
Stability	Biological function lost beyond 43°C		
Dissolution	100%		
Bioavailability	100%		
Dosage Form	Powder		
Route of Administration	Oral		
Pharmacokinetics			
The whole powder is dissolved in potable water at room temperature and drunk all at once. It is absorbed through the villi in the gastrointestinal tract and presented into the blood stream. It is progressively metabolized in the liver and then excreted renally.			
Pharmacodynamics			
Once in the blood, it is circulated to various tissues and organs. It never penetrates individual cells but remains within the circulatory system. It lyses cancerous cells, exposing their altered genetic material to normal biochemical pathways that eliminate them, which stops proliferation of cancerous cells.			
In Vitro Interaction of API with Some of the Common Drugs			
Category	Generic/ Trade Name	Effect on API Potency	
		Inhibitory	None
Antibiotic/ Antimicrobial	Amoxicillin		✓
	Ampicillin		✓
	Ampiclox (Ampicillin and Cloxacillin)		✓
	Ciprofloxacin	✓	
	Metronidazole		✓
	Erythromycin		✓
	Fluconazole		✓
	Seprin		✓
	Acyclovir		✓
Antimalarial/ Antiparasitic	Fansida		✓
	Quinine	✓	
	Artemether-Lumefantrine		✓
	Albendazole		✓
Analgesic	Paracetamol		✓
	Paracetamol Caffeine Aspirin	✓	
	Diclofenac		✓
	Ibuprofen		✓

	Tramadol		✓
	Piroxicam		✓
Antihypertensive	Nifedipine		✓
	Losartan		✓
	Atenolol		✓
	Furosemide (Lasix)		✓
	Propranolol	✓	
Antidiabetic	Metformin		✓
	Glibenclamide		✓
Antiasthmatic	Aminophylline		✓
	Salbutamol		✓
	Prednisolone		✓
Anti-allergic	Cetirizine	✓	
	Cimetidine		✓
Antihyperlipidaemic	Atorvastatin		✓
Anticoagulant	Cardiac Aspirin		✓
Minerals	Magnesium/ Aluminium		✓
	Zinc	✓	
	Vitamin C		✓
	Vitamin B ₁ , B ₂ , B ₆ and Niacinamide		✓

18-Jul-21

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Comment;
BKG
HIV1/2
HEPATITIS B
CANCER
COVID-19

ANALYSED BY:

Robert Mijumbi

18/07/2021

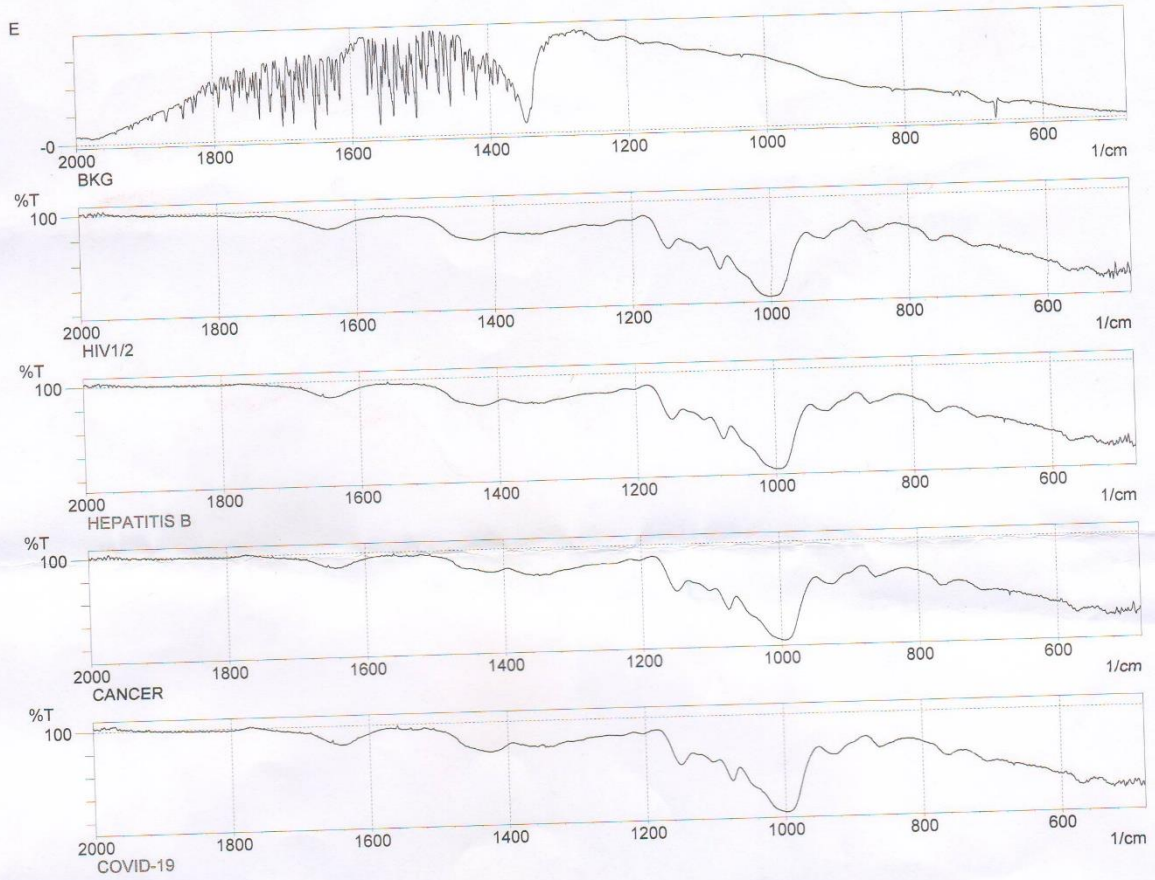
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Ajyka Pius

18/07/2021.

06-Aug-21

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Comment;
BKG
HIV1/2
HEPATITIS B
CANCER
COVID-19

ANALYSED BY:

Robert Mijumbi

06/08/2021

CHECKED BY:

Ojuka Pius

06/08/2021