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Spaces of argumentation and their interaction -Some elements of thought inspired by controversies and dispute in France during the Covid-19 crisis

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Motivation

- Work in progress, no formalisation yet;
- Covid-19: circulation of arguments in public spaces;
- Particularity: presence of scientists on the media;
- February 2020: scientific knowledge on SARS-CoV-2 virus constructed and shared in "real time" => emergence of *controversies*;
- Collision between media (short temporality) and science (slow and consensus not always synonym of truth);
- Interested in the interactions and exchanges between these worlds;
- In Zhe Yu and Shier Ju' talk: context is different culture background, norms and values and preferences over them. Change of a context: a strategy to get consensus. All participants share same norms (possibly with different preferences);
- Instead, our interest is at *debate switch* leading to opinion polarisation or incommunicability;
- How can different groups debate when sharing information, evidence, refer to different notions of acceptable arguments and proof standards?

Introduction

- March 2020: Dr Raoult learns by a Chinese colleague that chloroquine could work on the SARS-CoV-2 virus in vitro;
- Few weeks later: hydroxychloroquine (HCQ) can reduce viral load (even more in combination with azytromycine (Az));
- Results were not accepted because preliminary;
- The debate in France became very heated;
- Recorded some arguments that circulated;
- We have <u>no</u> medical competence to assess the correctness of the arguments;
- Our interest: exchange of arguments in a real case, between <u>different</u> worlds, with different notions of "rationality".



#FauciLeaks: 25/3/2020 email exchange between Dr. Anthony Fauci, American immunologist, who was the main adviser of Donald Trump and then of Joe Biden, and Jean-François Delfraissy, President of the Scientific Council:

On the therapeutic level, Yazdan Yazdanpanah has taken my place and who is now leading REACTing.

Tomorrow morning a french-european medical trial will begin, under the control of the WHO, with 5 arms (placebo/kalatra/kalatra +interferon béta / product of Guilead / hydroxychloroquine alone or in combination for next week).

As you may know, we are currently facing a press buzz since the announcement made by Dr. Raoult about the effectiveness of hydroxy-chloroquine. His data is not particularly convincing. We can distinguish a slight positive signal but it must be confirmed by a well made <u>randomised trial</u>. We also started a monkey-model study with HC and we should have the results by the end of next week. Also a cohort of severe COVID+ patients have been created and some will be treated with HC : we will analyse those results with all the consciousness needed. It will also be possible to do a prevention trial for healthcare

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professionals and aging people with HC. I have an enormous political pressure to release HC and to give it to everyone but I am currently resisting...

How is the situation in the USA, especially after Trump's announcement ? What is the NIH position ? What will be put into force ?

Thanks for your answer.

I put Yazdan Yazdanpanah in a copy of this email.

Best regard

Pr. Jean-François Delfraissy

Président | *President* Comité consultatif national d'éthique pour les sciences de la vie et de la santé **SHORT REVIEW**

Randomized controlled trials—a critical re-appraisal

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Abstract

Randomized controlled trials (RCTs) are considered to represent the gold standard of scientific studies and paved the way for evidencebased medicine (EBM). Besides the initial aim to improve the quality of patient care, EBM is used in the meanwhile for political and economic decision-making and legal issues as well. A review of the literature was performed, followed by a search using links and references of the detected articles. Additionally, homepages for German institutions of public health were screeened. Substantial limitations of RCTs and EBM health care could be identified. Based on the selected literature, 80% of the medical treatments have low evidence. RCTs are expensive and are mainly performed by the industry nowadays. A publication bias for positive results exists. Some RCTs are of low external validity. Many studies have a low fragility index. Nonetheless, negative RCTs could be of benefit for the patients. The results of RCTs, gained in a distinct patient population, are partially generalized. RCTs should be analyzed critically before adopting the results to daily clinical routine. It is not really justified to use RCTs and EBM for political and economic decisionmaking and legal issues as seen today.

Keywords Randomized controlled trial \cdot RCT \cdot Fragility index \cdot Evidenced-based medicine \cdot EBM



"My team and I believe we have found a treatment. And in terms of medical ethics, I feel that I have no right as a doctor not to use the only treatment that has been proven so far." (Interview D. Raoult, *Le Parisien*, 22/03/2020)



"To those who say that thirty multicenter studies and one thousand patients are needed, I answer that if we were to apply the rules of the current methodologists, we would have to redo a study on the interest of the parachute. Take 100 people, half with parachutes and half without, and count the deaths at the end to see which is more effective." (Interview D. Raoult, *Le Parisien*, 22/03/2020)



The New York Times Magazine

He Was a Science Star. Then He Promoted a Questionable Cure for Covid-19.

The man behind Trump's favorite unproven treatment has made a great career assailing orthodoxy. His claim of a 100 percent cure rate shocked scientists around the world.







Dr Gaetan Burgio, MD, PhD.

@GaetanBurgio

Huge observational study that just came out in @lancet on 96,032 #COVID19 patients including 14,888 treated with #hydroxychroloquine HCQ or CQ \pm AZ showing no benefit against #COVID19 but significant increase of serious adverse cardiac effects (QTc)

thelancet.com/lancet/article...





Les dernières études publiées sur l'hydroxychloroquine montrent une discordance entre les données observationnelles et les analyses rétrospectives de bases de dossiers de patients.

A l'IHU, nous faisons confiance à la réalité, pas au big data mal maitrisé



4000 patients traités VS Big Data : qui croire ? Bulletin d'information scientifique de l'IHU - Nous avons le droit d'être intelligents !Pr Didier Raoult, ... S youtube.com

12:25 PM · May 25, 2020

See the latest COVID-19 information on Twitter 12.1K 45



