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# Medical Anthropology Quarterly

# Multimorbidity, Polyiatrogenesis, and COVID-19

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## Multimorbidity, Polyiatrogenesis, and COVID-19

#### **Abstract**

To date, the strongest predictor for dying with COVID-19 is suffering from several chronic disorders prior to the viral infection. Pre-existing multimorbidity is highly correlated with socioeconomic inequality. In turn, having several chronic conditions is closely linked to multiple medication intake, especially in richer countries with good access to biomedical care. Owing to its vertical structure, biomedicine often risks giving multiple treatments in an uncoordinated way. Such lack of integrated care can create complex forms of iatrogenic harm. Multimorbidity is often exacerbated by a pharmaceuticalization of social deprivation in place of integrated care. In this article, I explore the possibility that clusters of overmedication are a contributing factor to higher death rates from COVID-19, especially in poorer areas within richer countries. Anthropological perspectives on the social embeddedness of multimorbidity and multiple medication use can expand our understanding of who is most vulnerable to SARS-CoV-2.

## "We really need to get data and we need to get data fast"

On March 18, 2020, Dr. Anthony Fauci and Dr. Howard Bauchner discussed a possible link between hypertension medications and a heightened risk of dying with a coronavirus infection. Bauchner is the Editor of JAMA, the Journal of the American Medical Association. Fauci directs the US National Institute of Allergy and Infectious Diseases. After joining the White House Coronavirus Task Force, Fauci became the world's most famous scientific advisor on COVID-19. In this conversation, Fauci highlighted possible links between ACE (angiotensin converting enzyme) inhibitors and COVID-19 fatalities. Fauci said that ACE inhibitors can increase "the expression of the receptors for the virus" (JN Learning 2020). Fauci was struck by reports from Italy that the vast majority of those who died with COVID-19 suffered from hypertension. Italy-then the European epicenter of the pandemic--was a rich country with excellent access to medical care, and many COVID-19 victims had been taking ACE inhibitors to treat their hypertension. "Why should someone who has hypertension that's well controlled have a much greater chance of dying than somebody else with any other kind of underlying condition?," Fauci asked. "We really need to get data and we need to get data fast" (JN Learning 2020). Fauci's concern is both about the clinic and the population. Significant patterns on a population level (here, Italian death rates) flag up patterns in clinical practice (widespread use of ACE inhibitors), which then lead to an exploration of pathogenesis (ACE inhibitors, cell receptors). I agree with Fauci that a careful

triangulation of pharmacological, clinical, epidemiological, and social science evidence is necessary to understand the action of the virus, how it spreads, and who is most vulnerable to it.

In the months following Fauci's call for data, a handful of medical articles appeared that examined the link between COVID-19 and antihypertensive drugs. The results were said to be inconclusive because none of the studies "adjusted for confounding variables" and so it remained unclear "if this association [between antihypertensives and high death rates] is related to the pathogenesis of hypertension or another associated comorbidity or treatment" (Patel & Verma 2020). The American Heart Association, a nonprofit organization that funds cardiological research and patient education, continued to recommend ACE inhibitors. In the absence of certainty about increased risk from ACE inhibitors, they told patients: "continue taking all your medications as prescribed, including ACE inhibitors ... If you are diagnosed with COVID-19, you should be fully evaluated before adding or removing any treatments ... your overall medical condition is much better if your blood pressure and diabetes are optimally controlled" (AHA 2020).

Fauci's call that "we need to get data fast" was only heard for ACE inhibitors but not for other medications, and possible links between the pathogenesis of COVID-19 and complex medication patterns remain entirely unknown. I argue that Fauci's challenge remains as important now as it was at the beginning of the pandemic. There are at least

two ways that chronic pharmaceutical use might put people at a higher risk of dying with COVID-19. One is that medications could directly augment virulence and pathogenesis (as intimated by Fauci). Another is that overtreatment adds to the overall morbidity index, regardless of any specific drug mechanism, thus making an overmedicated patient less healthy and less able to resist succumbing to the disease. In what follows, I will explore the possibility that multiple medication intake may lead to poor outcomes for coronavirus by this pathway. My goal is to show how medical anthropology, especially an anthropology of pharmaceuticals, may add a less obvious path in explaining the global impact of COVID-19 than that taken by virologists or epidemiologists.

While the pandemic modelling first anticipated that the coronavirus would behave like influenza, researchers realized that COVID-19 is a "complex multi-system clinical syndrome" (Roberts et al. 2020) that has a completely different trajectory than influenza. According to clinical reports as of this writing, the virus starts to spread in the lungs but can go on to attack all other organs, including the brain. Even when a patient survives the virus infection, the long-term effects, such as cognitive impairment, psychological distress, or reduced lung capacity, can be severe and protracted. Some of the long-term harm may even be caused by "iatrogenic damage" (Roberts et al. 2020), which comes from clinical uncertainty over how to best treat those with severe COVID-19 symptoms. It is now better understood that treating COVID-19 as if it was a form of pneumonia, e.g.

by connecting all patients with low oxygen levels to ventilators, may do more harm than good (Marini and Gattinoni 2020). The pathogenesis I will explore here is the one that connects victims' multiple chronic health problems to the risks of dying with COVID-19 specifically because of victims' multiple chronic medication histories. Could it be that socioeconomic deprivation coupled with chronic medication consumption is a predictor of dying with a SARS-CoV-2 infection?

To date, one medical study has appeared that shows polypharmacy being strongly correlated with the risk of *catching* Sars-CoV-2. In an analysis of half a million UK biobank users, McQueenie and colleagues found "that there was a clear dose relationship whereby the risk of a COVID-19 positive test rose steadily with polypharmacy level" (2020: 15). This study did not identify which kinds of medications were used, and it only explored infection risks.

My approach picks up on these threads from the medical literature to explore social factors linked to the spread and impact of the pandemic. First, I explore the social causes of multimorbidity and overmedication in relation to socioeconomic deprivation in one of the UK's most deprived areas where I recently conducted fieldwork. I then offer case studies that demonstrate how multiple medication use exacerbates patients' health. In the final section I explore population-level links between chronic illness, chronic medication use, and dying from an acute coronavirus infection. My suggestion is that medication patterns should be considered in studies on pathogenesis and epidemiology.

COVID-19 death rates seem to be highest (1) where a country has high rates of multimorbidity; (2) where multimorbidity is deepened by high levels of social inequality; (3) where the health system enables easy access to multiple medications; and (4) where multiple chronic treatments are insufficiently integrated. In wealthy countries, COVID-19 appears to claim most victims in poorer pockets *within* richer regions, and complex medication regimes might be a missing piece in this puzzle. Anthropological perspectives might offer novel ways of conceptualizing not just the social but also the epidemiological contours of the pandemic as problems of "local biologies" (Lock and Nguyen 2018).

# **Uncovering multimorbidity**

The clinical and epidemiological findings on COVID-19 and multimorbidity are, by now, fairly robust. From early on, medical reports found strong links between underlying health conditions and COVID-19. The majority of people infected with SARS-CoV-2, by some estimates 78%, remain asymptomatic (Day 2020). A minority of infected people experience mild to moderate flu symptoms. In perhaps 1% of people infected, COVID-19 turns fatal. Whether a SARS-CoV-2 infection kills seems highly correlated on the presence of pre-existing problems. The origins and forms of multimorbidity are thus extremely important for understanding global COVID-19 mortality patterns.

Already the first studies of mortality in China's Wuhan city showed that almost all the people who died with the virus had pre-existing disorders (Novel Coronavirus Response Team 2020). The data from China highlighted heart disease, diabetes, chronic respiratory disease, high blood pressure, and cancer (Novel Coronavirus Response Team 2020). Studies from around the world confirmed these findings. A study among patients admitted to New York City hospitals (Richardson, Hirsch & Narasimhan 2020) found similar conditions, especially hypertension (56.6% of COVID-19 patients), obesity (41.7%), and diabetes (33.8%). 21% of the COVID-19 patients admitted to NYC hospitals died. Using an assessment tool called the Charlson Comorbidity Index (CCI), which predicts 10-year survival rates in patients with multiple chronic problems, this study found that among all the patients admitted to New York City hospitals with a SARS-CoV-2 infection, half were expected to die of existing morbidities within the next 10 years.

Forensic pathologists in the German city of Hamburg performed full autopsies on hundreds of COVID-19 victims. In *every* case they found one or more of these preexisting conditions: heart disease, asthma, chronic obstructive pulmonary disease, peripheral artery disease, diabetes, obesity, or a neurodegenerative disease (Wichmann, Sperhake, Lütgehetmann et al. 2020). The Hamburg study discovered at least one preexisting condition in every patient examined, and multimorbidity in the majority of cases. The findings from Hamburg were widely reported in German media. Dr. Klaus

Püschel, the Director of the Hamburg Institute for Forensic Medicine, argued that SARS-CoV-2 is not nearly as serious a threat to the general population as it was believed to be. No one dies *only* of a SARS-CoV-2 infection, he said. At least one underlying health problem is always present (Wunder 2020).

Multimorbidity occurs when the same person suffers from two or more chronic disorders, which can be noncommunicable, infectious, or mental (Academy of Medical Sciences 2018: 6). There is no agreed way of how to measure multimorbidity: some researchers classify multimorbidity by how many disorders occur together, while others look for systematic clusters among co-occuring problems. What problems are counted in one of these clusters also varies, some researchers consider only a handful of chronic disorders as constitutive of multimorbidity (Dugravot et al. 2020), while others capture dozens of conditions (Payne 2020). Multimorbidity overlaps with other constructs, such as "frailty" (Hanlon et al. 2018) and "disability" (Stineman 2012).

The notion of "comorbidity," as the coexistence of an "index" disorder and a secondary disorder, emerged in the 1970s (Weaver, Barrett & Nichter 2016). Research into *multi*morbidity as a more complex category of diagnosis was almost inexistent a decade ago, but took off when the epidemic spread of co-occurring chronic disorders came into view (Busija et al. 2019: 1). Rising rates of multimorbidity have become a concern for both clinicians and epidemiologists. In rich countries, multimorbidity makes up 25-50% of the overall disease burden (Garin et al. 2016; van der Aa et al. 2017). In the

UK, for example, multimorbid conditions account for 50% of GP appointments, 64% of outpatient appointments, 70% of inpatient bed days, and 70% of total health expenditures (Aiden 2018: 4). Longer lifespans mean more multimorbidity: the older people are, the more chronic health problems they have. Up to two thirds of people over 65 are multimorbid. Treating older patients accounts for the majority of all health expenditures (Kaufman 2015).

Multimorbidity is strongly correlated with protracted medication uses. About half of older adults in richer countries are on five or more medications (Mangin & Garfinkel 2019: 1). The pharmaceutical industry promotes the regular consumption of five or more medications as necessary for the maintenance of health. For the industry, chronic polypharmacy is recommended as the new normal because multimorbidity is now normal. Already a decade ago, people in richer countries were on nine to thirteen prescription drugs in any given year (Dumit 2012: 2). Regimes of chronic polypharmacy originated in the US in the 2000s but have become normalized across the world wherever drugs are affordable and accessible.

Multimorbidity is not a new condition, as there have always been people with more than two health issues at the same time. But the growth of multimorbidity diagnoses and the biomedical focus on it are new. Dr. Chris Whitty, the UK government's chief advisor on COVID-19, argued in January 2020--just when the pandemic hit--that multimorbidity did not come into view for so long because biomedicine is organized "vertically" along

specific diseases, while a "horizontal" understanding of simultaneous disorders is lacking (Whitty et al. 2020: 1). Verticality increases the risk of iatrogenesis by way of multiple uncoordinated treatments. As others have shown, this verticality is itself an artefact of reliance on a biomedical model that tries to isolate causes of disorders and to develop therapies that are reductionistic (White 2006: 141-142). Although multimorbidity is a recognized and proliferating problem, research on the interactions between multimorbidity and protracted medication use is still largely absent.

Current biomedical research on multimorbidity could be of use in mapping possible research avenues in relation to Covid-19. Some of this explores nonrandom clusters of disorders (Academy of Medical Sciences 2018: 5; Busjia 2019). Other research explores how socioeconomic inequality congeals and expands chronic disease clusters. The mostcited study on multimorbidity is by Barnett and colleagues (2012). In this study, Scottish patient health records were scrutinized, and showed a strong correlation between socioeconomic deprivation and multimorbid suffering. A decade ago, already 23.2% of all Scottish patients were multimorbid and a third of these multimorbid patients had both mental and physical problems. Socioeconomic deprivation shifted the onset of multimorbidity forward: "young and middle-aged adults living in the most deprived areas had rates of multimorbidity equivalent to those aged 10-15 older in the most affluent areas" (2012: 39). In deprived neighborhoods, people in their 40s and 50s are as likely to have multiple chronic problems as 60 or 70-year olds in the average population. Barnett's (2012) quantitative results tally with qualitative studies on GPs' "endless struggle" with multimorbidity in deprived areas (O'Brien et al. 2011; Lawson et al. 2013). All available data suggest that multimorbidity keeps rising.

Even when multimorbidity is recognized by clinicians, there are few therapeutic protocols to deal with it (Guthrie et al. 2012). Given the logic of pharmaceutical industries, multimorbidity will continually escalate the number of pills required for treating the average patient. The more a health system is compartmentalized by specialization, the more likely it is that multimorbidity goes unrecognized as a problem of its own. Patients with multiple chronic problems experience biomedical care as lacking coherence (Schiøtz, Høst & Frølich 2016).

Some of the most important work in anthropology on multimorbidity is on "syndemics," which are "aggregation[s] of two or more diseases or other health conditions in a population in which there is some level of deleterious biological or behaviour interface that exacerbates the negative health effects of any or all of the diseases involved" (Singer et al. 2017: 941). The syndemics approach "move[s] beyond common medical conceptualisations of comorbidity and multimorbidity" (ibid.) by emphasizing structural violence (Farmer 1996) and social suffering (Kleinman, Das & Lock 1997). Syndemics researchers identify a number of multimorbid cluster patterns. For example, Mendenhall (2012; 2016) describes a pattern of "violence, immigration, depression, type 2 diabetes, and abuse" (VIDDA) among Mexican immigrant women

living in the US. "Immigration" tears families apart and leads to social isolation.

"Depression" is conditioned by traumatic childhood experiences and diminished selfefficacy. "Diabetes" is co-constituted by gender roles and food preferences. "Structural
violence" includes living in unsafe neighborhoods. Limited access to healthcare is
regarded as another form of structural violence.

A syndemics perspective helps to connect COVID-19 as "complex multi-system clinical syndrome" to multimorbidity, polyiatrogenesis, and social deprivation. In the next section, I will describe how people in a poor area of the UK experience multimorbidity and polyiatrogenesis, pointing to an as yet little explored syndemic pattern and also to possible ways we might rethink social vulnerability to SARS-CoV-2.

# Overtreating the underprivileged

My research took place in one of the UK's most deprived areas, which I will call Greyfield. Similar to the pattern described by Mendenhall (2012; 2016), Greyfielders experience clusters of structural violence, depression, diabetes, and abuse. However, they were not immigrants and they had universal access to healthcare, provided free of charge by the National Health Service (NHS). Thus, even the poorest are frequently on multiple medications without any financial burdens on households.

The population of Greyfield is largely white British. Most of the housing is made up of apartments in tower blocks and low-rises built in the 1960s and 1970s. The area saw

substantial redevelopments in the 1990s and 2000s. Despite some improvements,

Greyfield still has some of the UK's worst scores for health, income, employment,

education, and crime. Life expectancy of area residents is shockingly low at only 61 years

on average, which is 20 years shorter than the average UK life expectancy. Greyfield's life

expectancy is below that of Zimbabwe.

One of the infrastructural improvements to Greyfield since the 2000s was the construction of Greyfield Care Centre (GCC), a joint facility of the NHS and the local council. GCC tries to take a more integrated approach by combining different medical specializations (primary care, cancer support, dentistry, midwifery) with social services. GCC also houses a clinic for Lifestyle, Nutrition, and Complementary Therapy (LNC). My fieldwork was conducted in the LNC section of GCC. I analyzed 80 case files, discussed 30 selected cases with therapists, and shadowed 40 consultations. Most of the patients get referred to LNC by GPs in GCC. The patients that the LNC practitioners see are the same as those treated by the GPs, but LNC consultations allow far more time for patients to talk about their lives, their diets, and their medication regimes. LNC's hourlong, open-ended consultations allowed patients' to explore complex problem in much greater depth than in the GP consultations, where patients were given no more than five or ten minutes. The average length of GP consultations in the NHS—nine minutes--is one of the lowest among OECD countries (Wardle 2019). In the NHS, patients' time with GPs is extremely rationed, and longer consultations with a doctor are unavailable.

In LNC, Greyfield patients presented with a wide range of problems, but a few constellations came up repeatedly. Five or more co-occurring problems was the norm. The majority had mental problems along with physical problems. Traumatic life events were common. Chronically difficult social circumstances were nearly always present. Among various physical complaints, digestive diseases were virtually always present. Obesity was very common and was usually coupled with diabetes. Hypertension was pervasive. Physical problems appeared together with mental problems. Depression and anxiety were the most common mental illnesses.

Both mental and physical problems were often related to life trauma. Many traumatic events happened in relation to family members. Domestic violence and emotional abuse in intimate partnerships were common. Loss of family members, often at a young age, was another frequent source of trauma in this population. Miscarriages were also common. Haunting memories of losses suffered were a recurrent problem. Susan, a woman in her early 50s, for instance, first came to LNC because she wanted to lose weight and sleep better. In the first consultation, she broke down in tears. The sleep disturbance had started the previous year, when her brother died and one of her surgeries went wrong. She said her family was "broken": mother, father, cousins, uncles had all died, and her twin sister died when she was still a child. She said she often wakes up at night in a panic, as if a bright light was shining into her eyes.

Substance use was mentioned often by patients. Mostly, this referred to alcohol (for patients of all ages) and cannabis (among younger patients). Feeling "empty" or "bored" were patients' explanations for why substances were used. Strong food aversions were also common. For example, Jack, a patient in his late 30s, complained about depression, panic attacks, acid reflux, sleep disturbance, and headaches. He also suffered from severe back pain, which two previous surgeries failed to make better. He traced his problems to his childhood: he was always disgusted by eating meat, but his parents forced it down his throat. In his experience, he said, the best remedy against the pain and the panic is smoking cannabis. Patients like Susan and Jack presented with complex multimorbidity in ways that are probably similar to many clinics in the world where a combination of poor diet, precarity, and multiple social traumas coalesce to cause or augment physical suffering.

Despite the common presentation of patients at Greyfield, and despite the ample access to care they were provided, their overall state of health seemed often worsened by uncoordinated treatments. Molly's experience exemplifies this vividly. Molly was a 37-year-old woman born and raised in Greyfield. Molly came to LNC for a persistent pain in her foot. In the first consultation with Heather, the LNC practitioner, Molly added that she was also suffering from diabetes type II, that she wanted to lose weight, and that she could help with her irritable bowel syndrome (IBS). Molly was aware of the main food triggers for her IBS but she could not keep it in check. Molly said that her job as a

care worker was very stressful, especially when working late hours. She said her stress "gets to my stomach." Her bowel movements were very irregular. The first time she was diagnosed with IBS was when she was a child. She got some treatment and the problem was not too troubling until five years ago when she had a protracted and difficult breakup with her husband, during which she and her son suffered domestic abuse. That led, in turn, to lots of "behavioral problems" in her son that she was still struggling with. Asked by Heather if she had time left to look after herself, Molly said "my self-care is really bad, I know that it is really bad."

Molly took a dose of Imodium every morning and had become completely dependent on it. Imodium (loperamide) is a common over-the-counter drug for acute diarrhea. The medication slows down the digestive process and should not be used for more than 48 hours. Molly complained about her daily eating habits, especially too many potato chips: "Pringles are my downfall." Molly drank three to four litres of Coca-Cola throughout the day: "I am really bad with fizzy drinks." Asked if the GPs had explained the links between diabetes, blood sugar and high-sugar drinks, Molly said that the doctors did not yet have time to talk about food. Instead, they were "just firing tablets at me." She had not been told by anyone face-to-face how to manage her diabetes. The LNC practitioner asked about Molly's blood sugar readings, but Molly replied that she had no idea. Heather fetched a blood sugar kit and administered a test, by pricking one of Molly's fingers and putting the blood in a glucose monitor. The reading showed

an extremely high glucose level of 17 mmol/L. Heather asked what medications Molly is on. Molly pulled a black rubbish bag out of her handbag and put it on the table: "my lucky bag of medicines," she laughed. Molly said that she is taking two medformin (a standard tablet for diabetes) every morning, plus another diabetes medication. Heather said that Molly should find out how to monitor her blood sugar levels, not least because her levels are high *despite* already being on drugs.

Molly unpacked seven other drugs from her "lucky" bag: an antidepressant (citalopram 20mg), propanonol (a beta-blocker, against the anxiety of having diarrhea while out of the house); Fianola, a contraceptive; an antihistamine for allergies; omeprazole against gastric reflux; gentamicin, an antibiotic drug; and a high-dose ibuprofen for the pain in the foot. Molly continued to unpack a number of alternative remedies, including sage pills and matcha green tea pills, which were meant to help her against "hot flashes." She said that she is a "hot person," "burning up very quickly," and joked that she should "move to Iceland" to feel more comfortable.

Heather asked about Molly's body weight. She said she weighs about 20 stone. At her body height, her BMI was 45, putting her into the obese category. Molly said she has always been "big." She then asked about a growth on her foot and showed it to Heather. A GP had told her it was a "ganglion." It caused her much pain and restricted her movement. By overcompensating with her other foot, she developed plantar fasciitis and dislocated a bone. She had phoned her GP about the ganglion but was told that there

was nothing else they could do. Heather probes further about what kind of pain Molly has in her foot and is told it is like "pins and needles," which Heather said might be related to the diabetes. Molly further said that she has not been told by any doctor what she could do about the plantar fasciitis. Drawing the consultation to a close, Heather said she does not want to prescribe any therapy yet. Instead, Molly should come back for a full assessment and then they would work out a care plan. Molly thanked Heather for taking so much time, saying that GPs only ever spoke to her briefly and only about a specific ailment. Heather went on to add that Molly should stop drinking fizzy drinks immediately. Molly asked if juices or fizzy water would be alright to have instead, but Heather told her that neither of them would be good for her, both for the sugar and for the fizz.

Molly's case gives an insight into the "invisible epidemic" (Mangin and Garfinkel 2019) of patients with multiple health problems receiving multiple poorly integrated treatments that cause too often cause more harm than good. My point is not that Molly's case is exceptional but rather that it is not exceptional at all. In fact, the cluster of multiple comorbidities experienced by Molly could be said to appear in syndemic form in many poor pockets of rich nations around the world. These pockets reveal high levels of multimorbidity and polyiatrogenesis. What I am suggesting is that such areas of social deprivation and overmedication might be at a substantially higher

risk from a virus that becomes particularly dangerous for people with multiple chronic conditions.

Molly's overflowing bag of medicines is an example of how multimorbidity is deepened by the use of too many pharmaceuticals. I am not arguing that any single one of the GPs' treatments is irrational. What I am arguing is that a vertical system does not give physicians enough time to control for potentially iatrogenic effects, especially in chronic patients. When multiple single-target medications are coupled with a lack of explaining what is going on, the risks of overtreated ills become even greater. In an era of rising multimorbidity, biomedical specificity becomes part of a new kind of pathology.

What is so striking about multimorbid patients in the UK is that their poor health cannot be attributed to a biomedical treatment gap, in the way that this might be done in Mendenhall's (2012) study on syndemics and structural violence in the US. The UK situation also differs from places like Brazil where poverty is heavily pharmaceuticalized (e.g., Scheper-Hughes 1992; Biehl 2005; Béhague 2015). In the UK, GPs are basically too short of time to treat complex patients in complex ways. They throw a pill at any ill in the belief that this constitutes good care while being unable to deal with the underlying problems and while losing sight of iatrogenesis. Greyfielders are deprived socioeconomically, but this deprivation does not translate into a lack of healthcare. The problem is a lack of integrated care.

## Pharmaceuticalized pockets within richer regions

It is still early to know the links between multimorbidity, polyiatrogenesis, and COVID-19. As mentioned, there is not a single medical study yet on links between COVID-19 fatalities and multiple chronic medication uses. No one is tracking systematically what kind of medications COVID-19 victims are consuming when they get into hospitals with severe symptoms, let alone how chronic medication regimes interact with COVID-19 in the community setting.

Even without this clinical evidence, it is possible to look for patterns on a population level. In the UK clinic I studied, high levels of multimorbidity and polyiatrogenesis were evident. If the hypothesis of chronic polypharmacy as a risk factor for COVID-19 holds, we would see higher rates of COVID-19 deaths in countries with high rates of multimorbidity, social inequality, ready access to biomedicine, and pronounced verticality in service delivery. What I found through ethnographic work in a UK clinic would lead to expect that death rates are indeed higher in the UK than in countries where this syndemic pattern is not as pronounced.

The global epidemiology of COVID-19 is still in flux, and many puzzles still have no convincing solution. Nevertheless, some epidemiological patterns have crystallized and remain fairly stable. First, we now know that the UK had become one of the world's hardest-hit countries by May 2020. While it is still difficult to draw meaningful

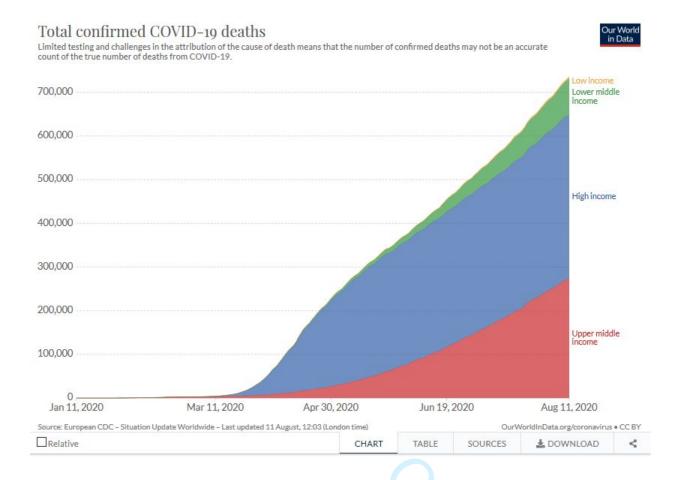
comparisons between countries, there is solid evidence that the UK was exceptionally affected by the pandemic. By August 2020, the UK had one of the world's highest COVID-19 death rates, both by absolute numbers of victims (more than 50,000) and by fatalities per million population (685 compared to 94 in the world) (Our World in Data 2020). The UK started to record "extremely high" excess death rates from week 13 onward (EuroMOMO 2020). While most other European countries saw their excess death rates return to normal levels over the next three weeks, and while other nations saw their case fatality rate decline as more people tested positive without deadly outcomes, the UK continued to record "extremely high" excess rates for another eight weeks. Even the UK's nation-wide lockdown took two months to bring a significant reduction in death rates. In June, a full twelve weeks into lockdown, the UK recorded daily death rates higher than all other EU countries combined. The UK has done worse than most countries in the world, including the US. There were abject errors in the UK government's COVID-19 response (Horton 2020), but these policy failures cannot explain the UK's exceptionally high death toll.

Studies on European death rates consistently found correlations between social deprivation and dying with the virus: "COVID-19 is painfully exposing the existing and persisting health inequalities in our societies. This pandemic will have the heaviest impact on the lives of people living in deprivation or facing difficult socio-economic circumstances" (EuroHealthNet 2020). The UK showed the same pattern. Across

England, Wales, and Scotland, death rates were more than twice as high in socioeconomically deprived areas than in other areas. People in low-paid employment were dying at much higher rates than those in white-collar employment (New Statesman 2020). Socioeconomic inequality is one of the strongest predictors for dying with COVID-19 (Clark & Whiteley 2020).

In the UK, as in practically all the European countries, social inequality does not translate into unequal access to biomedical care. In Greyfield and elsewhere in the UK, relative poverty does not mean less access. But no amount of clinical care can shield from dying with COVID-19. Docherty and colleagues' (2020) study on hospital outcomes in the UK found that the death rate of all patients admitted was 33%. In ICU units, 45% died. The UK is not an outlier in poor outcomes of hospital care. Reflecting on her gruelling experiences of giving intensive care to COVID-19 patients in New York City hospitals, Dr. Helen Ouyang (2020) realized that all the biomedical care in the world was unable to save people's lives: "What I think will actually cause moral injury is seeing people die after getting the most advanced care available. People who come in talking, with stories to share. They get care—the best that modern medicine has to offer—with life-prolonging machines and IV drips of all sorts of critical-care drugs. We put our full minds and whole hearts into trying to save them. Then I see their bodies shut down anyway." Even the best-equipped healthcare system offers no good defense against COVID-19.

Compared to other infectious diseases, COVID-19 behaves in an utterly surprising way. All other major infectious diseases—tuberculosis, HIV, or malaria—show a strong correlation between low GDP and high death rates. SARS-CoV-2 inverses this pattern on: so far, the impact of the virus was far more severe in rich countries than in lowincome countries. To be sure, the pandemic is still unfolding and what the situation might be in the long run is not known. And yet, to date high GDP and high COVID-19 death rates are strongly correlated. By mid-August 2020, upper-middle and high-income countries suffered 90% of all COVID-19 deaths, despite having about half the world's population. This pattern is so strange because infectious diseases usually strike much iauso harder in poorer countries.



Until July 2020, all the world's countries with the highest per million population death rates lay in a corridor stretching from Northwest to Southwest Europe. Sweden, a country famous for not implementing a lockdown, only joined this group in the last days of May. Death rates per million population of 500 or more outside these European countries have so far only been reached in the US, Chile, and Peru, and that only since July. Viral infections usually take advantage of "multiple co-infections and biological and social vulnerabilities in a world where syndemics disproportionately affect marginalized

and less-resourced communities" (Manderson & Wahlberg 2020: 2). That means, practically without exception, that infectious diseases are far worse in poorer countries than in richer countries. COVID-19 turns this pattern on its head: all the hardest-hit countries are GDP-rich countries. COVID-19 does not seem to strike most where levels of absolute poverty are high, instead it hits the rich countries. There were some outliers, notably East Asian countries did far better than what their GDP per capita would lead to expect. The overall trend is clear, however. The world map of COVID-19 mortality does not show any Global North/South distributions along poor access to quality healthcare. Instead, the COVID-19 map looks like an atlas of rich industrialized countries with excellent clinical care. To be sure, GDP is a poor proxy for ease of access to biomedical treatments—US healthcare is a case in point. At the very least it safe to say that high income and excellent biomedical care do not protect against high COVID-19 death rates.

The country-to-country comparison hides that *relative* deprivation within a wealthy context is a key risk factor for health (Ecks 2021). COVID-19 strikes poorer pockets *within* richer regions. Relatively deprived communities do not necessarily lack access to biomedical care. These communities may have high levels of multimorbidity *within* a context of relatively good healthcare access. Good access to healthcare clearly does not protect anyone from dying with COVID-19. In fact, easy access may translate into chronic overmedicalization, which in turn may make people more vulnerable to dying with the virus infection. The UK data on double mortality rates in poorer areas confirm

this pattern. Other countries with stark regional and local wealth disparities exhibit this pattern as well. Relative deprivation leads to higher rates of multimorbidity.

Multimorbidity increases the risk of polyiatrogenesis, especially where treatments are accessible and affordable. It is possible that COVID-19 particularly affects multimorbid patients who are prematurely sick because of deep wealth inequalities and whose multimorbidity is exacerbated by polyiatrogenesis.

It is possible that there are other factors at work that explain why GDP-rich countries have much higher death rates. Some of them are about GDP and global connectedness; others are about GDP and population age structure. SARS-CoV-2 probably first made impact along global air traffic routes, so that many poorer countries might have had a delayed exposure to the virus. But date of exposure seems unrelated to eventual death rates. Thailand, for example, was the first country outside China to record a SARS-CoV-2 infection, on January 13, 2020. The country has a population of 69 million. By August, it had 58 deaths. The UK population is 66 million. The country recorded its first SARS-CoV-2 case on 28 February 2020, seven weeks after Thailand. By August, the UK had more than 50,000 deaths. Per one fatality in Thailand there were 862 fatalities in the UK. Well into June 2020, the UK recorded five times more daily deaths than Thailand's *entire* death count in a population larger than the UK's.

Many argue that richer countries, especially in North-Southwest Europe, became the global epicenter of the pandemic because people there have longer life expectancies and,

therefore, higher rates of multimorbidity. But longer life expectancies alone cannot explain any of these patterns. Among the Europeans, only Italians and Spaniards are in the top ten globally. The UK ranks only 29th in the world for life expectancy. Japan, Hong Kong and many other countries have much longer life expectancies than the UK, but are not nearly as much hit by the pandemic. The same point can be made about a population's median age as a protective factor against high COVID-19 death rates. That low-income countries have younger populations and "therefore" have lower COVID-19 death rates is taken as an obvious fact (e.g., Pulla 2020). But if median age were a vital factor, why would Peru have a COVID-19 death rate of 651 per million population at a median age of just 27.5 years? At 46.3 years, Japan has the world's highest median age; at 8 deaths per million population, Japan also has one of the world's lowest COVID-19 death rates. Neither a country's life expectancy nor its median age seem to be decisive factors in COVID-19 death rates.

To date, none of the available epidemiological and population data can confirm or disconfirm links between high rates of chronic medication uses and COVID-19 death rates. High GDP alone does not translate directly into high rates of pharmaceutical consumption, fragmented care, or deeper wealth disparities within countries. But the fact remains that the risk of dying with COVID-19 is not correlated with low income levels, as practically all other infectious diseases are. Nor is it related to poor healthcare access, as is the case with all other infectious diseases. Instead, epidemiology of COVID-

19 seems to confirm that countries with high rates of multimorbidity, high levels of internal social inequality, and good health access are hardest hit.

Checking COVID-19 victims not just for underlying health conditions but also for medication histories is the only way of knowing if there might be a link to chronic multiple medication use. Much more research is needed to figure out how susceptibility to dying with COVID-19 might be related to medication patterns. But to date, neither clinicians, epidemiologists, nor social scientists are studying this. We need to link COVID-19 deaths to clustered disorders and we need to link clustered disorders to clustered overmedication. The short ethnographic case study I presented here is only a tiny fragment in the work that needs to be done on experiences and causes of multimorbidity, chronic medication use, and polyiatrogenesis in a pandemic era. Both clinical and epidemiological findings are suggestive of links between relative socioeconomic deprivation, medication overuse, and susceptibility to the novel coronavirus. As long as there is no effective medication to treat COVID-19, clinicians should make sure that existing treatments for other medical conditions are used with greater care.

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