The recent changes to covid-19 restrictions indicate a new phase of the pandemic, where we are supposedly learning to live with the virus. Masks are not required, testing is no longer readily available to the public, and isolation has become optional rather than mandatory.

I’m sure that many businesses are pleased to see an end to testing and isolation, in the hope that covid-19 related absences among their workforce will now rapidly decline. But what about the NHS? Last month the remaining population controls against covid-19 ended. As a result, cases rose steeply throughout the population, leading to high absence rates among NHS staff as they were off sick or needed to self-isolate.

This has inevitably meant cancelling operations, outpatient appointments, and diagnostic tests. We’re still declaring covid outbreaks on many of our wards—leading to fewer available hospital beds, which has a further detrimental impact on waiting lists and patient flow through the NHS.

When the message being relayed to the public is that the pandemic is virtually over and that the NHS is in some sort of accelerated recovery mode, it’s no wonder that patients have a low tolerance for more news of cancellations. It’s adding fuel to the fire when public satisfaction with the NHS is already at its lowest recorded level since 1997.

Last week NHS England announced that infection and prevention control measures introduced during the pandemic were being relaxed, partly to help free up capacity in healthcare services to tackle the huge treatment backlog. I’m afraid that, no matter what infection prevention and control measures we take in the workplace, we’ll inevitably still be exposed to the virus in our lives outside work. This is crippling our ability to do the much needed catch-up work now facing almost every sector of the health service.

Worn-out NHS workers are the ones caught in the middle of this bind—either through being acutely unwell with covid and having to take time off or working in understaffed departments, unable to provide a decent service. I dread to think of the impact of long covid on top of this, as it’s yet to be fully felt or at least captured in available statistics—but recent data from the Office for National Statistics warn that many of the people affected will probably have symptoms for a year or more.

Suffice it to say, this is not the recovery that any of us had anticipated or hoped for. With a stretched, depleted, and unwell workforce, the NHS’s staff and its patients have tough times ahead.

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This is not the recovery that any of us had anticipated or hoped for.
How trainees can find lessons in clinical administration

What feel like mundane tasks are the grease on the NHS’s wheels

As I’m the most junior doctor in the acute medical unit, each of my days starts in a similar way. I take a piece of off-white NHS paper, which has boxes and bed numbers already printed on it, and add patients’ names down the side. During the board round, I keep a careful track of the tasks that need to be completed for each patient.

I use the classic box system—which is somehow passed down through generations of medics without anyone ever really explaining to you how it works—to keep track of my progress. Typical jobs include filling in referral forms, negotiating investigations with radiologists and endoscopists, taking blood and placing cannulae, and gathering collateral histories from relatives.

My day-to-day work mostly falls into the category of “admin.” A frequent complaint of junior doctors, especially those in foundation year 1 (FY1) who don’t have full GMC registration, is that too much time is spent doing these tasks, rather than being trained through clinical contact. Indeed, this is a longstanding international concern, with Forbes reporting in 2016 that doctors “waste two-thirds of their time doing paperwork.”

I agree there is always a conversation to be had about ensuring training is fit for purpose and that it is allowing juniors to develop the skills they need to progress. Yet in reflecting on my first job as a junior doctor, I see these seemingly small “admin” tasks differently: as an opportunity to learn about myself and my approach to a clinical workload in a relatively controlled and supported environment.

Each morning, it’s up to me to prioritise the jobs I do. While senior doctors might emphasise certain tasks that need to be completed as a matter of urgency, I am responsible for how I conduct my working day to maximise the chances of the patients who are under my care being seen by the right people and investigated in the right way.

Managing workload
Gaining experience in how to prioritise these “low stakes” (that is, not immediately life or death) tasks gives junior doctors an opportunity to tentatively begin to manage their clinical workload.

I have found that the more experience I have navigating the complexities of these daytime jobs, the more confident I felt in prioritising my time during on-call shifts, where there was often a higher level of clinical prioritisation required.

I have also come to value the learning I’ve gained from communicating with other members of the multidisciplinary team during these “admin” jobs. Take the quintessential task of phoning the on-call radiologist to discuss a request for a scan. You feel under great pressure to secure this scan and are nervous. But after the first few times, you learn to anticipate the questions you may be asked, and eventually come to the conversation armed with all the information you might need to secure the scan.

These interactions not only teach us about professional communication—they allow us as very junior doctors to explore our clinical “personalities” in how we approach interactions that have the potential to be challenging. Ultimately, this experience of clinical communication is valuable; when it comes to discussing important clinical handovers—say, about a deteriorating patient on a night shift—you are able to go into the

Rising costs spell disaster for the nation’s health

Black storm clouds are hanging over the nation as inflation rockets, national insurance rises, and the energy price cap lifts—unleashing escalating energy prices across the UK.

This perfect storm of rising costs will disproportionately affect the health of the poorest in society, adding to already unacceptable wide inequalities in the UK.

Energy bills are expected to rise by almost £700 this year for a typical household. Shockingly, this means that energy bills are expected to rise by more than 50%. This is five times the previous biggest increase of £139, which came into effect only in October last year. As a result, more than one in four households in the UK will be living in fuel poverty.

Fuel poverty is part of a cycle of poor health because of the many interactions between

It is likely a further 1.3 million people will be pushed into absolute poverty challenging socioeconomic circumstances and societal inequalities. Living in cold temperatures increases the risk of respiratory infections and exacerbates many chronic conditions. The indirect effects of fuel poverty are also concerning; households that spend a greater proportion of their income on energy have less to spend on food and other health necessities. Financial insecurity drives greater social isolation, and carrying a greater burden of debt causes mental health problems for both adults and children.

Politicians may say that we must bear the rising cost of living together, but the problem is that these costs are not equally distributed. People who live in low income areas do not have lower fuel bills. They also bear the brunt
conversation with the confidence that you have all the information you might need.

In my experience, your FY1 year is not simply a rite of passage or a job to be done because “someone has to do it, and you only have to do it for a year” (as one of my consultants so succinctly put it). These admin tasks, so often the domain of the most junior member of the team, are the grease on the wheels of the NHS, and they have a huge bearing on a patient’s journey and experience.

Alongside this, managing your own workload of these sometimes seemingly disparate, isolated tasks sets the blueprint for how you approach a clinical workload throughout your career; communicate with other healthcare professionals, patients, and relatives; and begin to lead a team. So to my fellow FY1s: I see your beautifully filled in referral forms, your deft handling of a tricky collateral history over the phone, or that time you got the radiologist to approve a scan that led to a diagnosis. Enjoy the small victories—you’re doing a great job.

Anna Harvey Blueme, academic foundation doctor, North East and North Cumbria Integrated Care System

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of decades of inaction in improving home insulation. It is the poorest who will struggle to keep afloat, and whose health will suffer, when the financial storm hits.

The UK government says it has done enough to support households through the cost of living crisis, but this support is not fairly weighted to protect the poorest in society.

The consequences of government policies and treasury decisions are that it is likely that a further 1.3 million people, including 500 000 children, will be pushed into absolute poverty over the next year. This will spell disaster for the health and wellbeing of the UK’s poorest citizens who already suffer substantially poorer health than their richer counterparts.

Neena Modi, president, BMA

Cite this as: BMJ 2022;377:o938

GP numbers have fallen, yet annual GP and practice nurse consultations have risen by 16%
Adjusting our expectations

I recently attended a workshop exploring the effects of switching to a “remote by default” model of general practice. This was made necessary for infection control at the height of the pandemic. The reasoning goes that if there’s no need to physically examine a patient to reach a diagnosis, why bother being in the same room?

In the real world, a complex mix of factors should influence our choice of consultation method, not all of which are considered when setting up new models of care. In many surgeries, remote consulting brought with it a triage system where appointments are allocated depending on information provided on an online form. This should enable practices to work out who really needs face-to-face GP interaction and where a phone call or help from another member of staff might be appropriate. Although always expressed in efficiency terms, this is often about rationing access to a scarce resource—one of a shrinking number of GPs.

Although offering some sort of e-consultation is now compulsory, some practices now require an online form for every appointment, while others bury the link on their website and carry on as before. In the workshop we were asked to consider scenarios where patients struggled to navigate systems. Some described patients’ frustration being interpreted as hostility by overburdened reception staff. In others the patients gave up, unable to fill in a computer form and lacking the time or phone credit to stay on hold for help to do so. The clear risk is that we miss a chance to cure, and the next presentation may be with a more serious illness or even an advanced malignancy.

There was consensus among the clinicians and patients present that most of the problems could be avoided (or ameliorated) by continuity of reception and clinical staff. However, I was disappointed to hear this was “not a realistic response.” Apparently, I have unrealistic expectations when I describe my version of what good general practice looks like: long term relationships, easy communication, and timely access. This is the service most people would like for themselves and their relatives. So, when did we agree to settle for less? In whose manifesto did it say that GPs would switch to an impersonal service reached by an online form, where your stated problem would be allocated to a doctor you didn’t know or your request for help might result in generic advice delivered electronically?

I’m encouraged by the committed young doctors joining our profession, but if we’re to keep them, and to stem early retirements, we need an injection of hope. We need to feel confident the things we value as GPs will be recognised and preserved—and that we, in turn, will be respected and listened to.

When did we agree to settle for less than long term relationships, easy communication, and timely access?

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Cite this as: BMJ 2022;377:o1106

Creativity and wellbeing

Paula Redmond, a clinical psychologist who supports healthcare workers, joins this podcast episode to talk about the common struggles NHS staff are going through right now, protective factors against burnout, and how creativity has helped her during difficult periods. She starts by describing the impact on healthcare professionals of not being able to deliver the care they want to: “It has an insidious, eroding effect on self-esteem and our connection with work—particularly for health professionals when we are so values driven.

We come into this work because we want to make a difference and help people, and often our identities and sense of self-worth are really tied up in that. But it means that when things are going wrong at work, when we’re not able to make a difference, it can really cut to the core of what’s important to us and how we feel about ourselves as people. Our professional and personal identities can get so fused that when one is struggling, it’s really hard to function in any other aspect of our lives.”

Redmond shares how being creative has been a source of joy and accomplishment for her:

“Creativity is a real, innate part of being human. Even if we’re not creating, we can surround ourselves with beauty and that’s good for us. Something like crafting, for example, combines both pleasure and mastery, which are two factors that are key in terms of lifting mood. If we think about some of the activities that we might do to help us relax—watching TV or scrolling on our phones, for example—we can spend hours doing that and we might feel more relaxed, but we don’t necessarily feel fulfilled. Whereas craft is something that can combine both the sense of being engaged in something beautiful, but also a sense of achieving something.”

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Edited by Kelly Brendel, deputy digital content editor, The BMJ
Can the world become a place where the planet and people flourish post-pandemic?

Covid-19 has impeded achievement of the sustainable development goals and a radical rethink of the global economy is required to meet them, argue Fran Baum and colleagues.

In 2015, the world adopted 17 sustainable development goals (SDGs) with 169 targets to be achieved by 2030. These goals aimed to create a world in which people and the planet flourish. They were more ambitious than the previous millennium development goals and linked human wellbeing with the sustainability of the planet. Achieving these goals would make the world fairer, more sustainable, biodiverse, and healthy as well more participatory, decolonised, and democratic. Yet even before the covid-19 pandemic concerns emerged about whether governments had the will to achieve these aspirational goals. Covid-19 has cast further doubt and seen reversals rather than progress on many of the goals.

We examine the effect of covid-19 on progress across the five inter-related dimensions of the SDGs—planet, people, prosperity, peace, and partnership—and discuss the political, social, and economic transformations required to meet them. Although creating new challenges for the SDGs, covid-19 has shown that governments can change policy rapidly when they want to. Similar rapid changes are now needed to advance progress towards achieving the SDGs, including radical reforms to fiscal and economic systems to reduce inequities and devise policies that confront the interests of elite groups.

**KEY MESSAGES**
- The UN sustainable development goals were intended to create a sustainable planet and a world in which all people could flourish by 2030
- The covid-19 pandemic has set back the achievement of the SDGs
- It has worsened social determinants of health, increased socioeconomic inequities, and restricted civil society activism
- New economic models are needed that enable strong social security, education, and health systems, and encourage participatory democracy

**Planet’s ability to support human life**

Planetary health and tenable human life on earth are at risk. Irreversible climate change will have a huge negative effect on health, particularly for those living in poor and marginalised settings. Unless there are immediate, rapid, and large reductions in greenhouse gas emissions, limiting warming to 1.5°C or even 2°C will be unachievable.

Although the pandemic has had some positive effects on the planet (eg, reducing air travel), some countries are reducing their environmental safeguards and seeing natural resources as “capital” on which to build their post-pandemic economic recovery. Countries rapidly adopted new policies to limit the human and economic effects of the covid-19 pandemic, yet no government has similarly acted for the arguably greater existential threat of climate change.

Nothing short of rapid, transformative change to protect ecosystems and reduce carbon dependence will be enough to safeguard planetary and human health. Our best guides to this future may be indigenous peoples around the world, who consider all living things to be connected and the planet’s best guides—livelihoods based on principles of living in harmony with nature and protecting the liveability of the planet.

**Indigenous peoples may be our best guides—livelihoods based on principles of living in harmony with nature and protecting the liveability of the planet**

Lives in which people can flourish

The SDGs envisaged a world in which people can flourish, where poverty and hunger are reduced, and extreme poverty is eliminated by 2030. Health and wellbeing, access to quality education, and the reduction of socioeconomic, gender, and other inequities were also seen as critical for people to fulfill their potential (table 1). Even before covid-19 it was projected that 670 million people would be extremely poor in 2030. The World Bank estimated that collapsing supply chains and economic contraction associated with the pandemic pushed around 97 million more into extreme poverty in 2021 raising the estimated total to 732 million people.

Disparities in covid-19 infections and mortality were experienced by minority ethnic groups in various parts of the world and reflect longstanding patterns of racism. The pandemic also contributed to increases in violence against women, girls, and LGBT+ people, and higher rates of poverty and food insecurity among women. Women experienced greater income and employment losses than men as well as increased domestic caring demands. Globally, the pandemic both revealed and exacerbated the social protection gap between high and low income countries, with health inequities likely to increase.

In his July 2020 report to the Human Rights Council, the special rapporteur
on extreme poverty and human rights noted that “poverty is a political choice” and eliminating it requires transformative policies including tax justice and redistribution, universal social security, and equitable participatory governance. The adoption of participatory governance is essential to hold governments to account and strengthen the will for such redistributive policies.

There is enough wealth in the world for all countries to meet their SDG targets; distribution, not scarcity, is the fundamental and policy amenable problem. Social security, public health, and education systems need to be based on the right of all people to have access to them as such systems are essential to redress systemic inequities that marginalise social groups and to ensure that quality healthcare, childcare, and education are accessible to all.

## Prosperity for all

The SDGs are also intended to provide the means by which all people can enjoy a prosperous and fulfilling life. An important pathway to prosperity is ensuring decent work through strengthened labour rights, protection of trade unions, and proper pay and working conditions. Social protection measures that safeguard people against shocks and stresses are also critical to decent work. Yet the pandemic has seen those already in informal or casual work being more likely to lose their jobs and at increased risk of covid-19.

Table 1 provides examples of how wealth inequalities have worsened markedly during the pandemic, with 2020 recording the steepest increase in global billionaires’ share of wealth on record. The 2022 World Inequality Report notes that the poorest half of the global population owns just 2% of total global wealth, while the richest 10% now own 76%. In wealthier countries, governments rapidly introduced massive income and business support programmes that saw public debt soar in a way that would have been unthinkable before the pandemic. This had a mixed effect: although the support maintained most people’s pre-pandemic income, it created “cheap money” for banks and investors in a liberalised and under-regulated financial system. Those owning companies directly benefiting from a pandemic (eg, drugs and online sales) or with the financial means to speculate more broadly (eg, in equity
markets, derivatives, real estate) saw their wealth increase substantially.9 At the same time millions of people around the world struggle for the prerequisites for healthy living. A small progressive tax (eg, 2%) on wealth could be used to improve access to health services, quality education, social security, and better work conditions for those with fewer resources.

A fundamental change is needed in the way prosperity is understood and measured. Achieving the SDGs will require rethinking the idea that an economic system must be based on GDP growth. Instead, alternative measures of prosperity need to be adopted, such as the genuine progress indicator12 or the happy planet index,13 both of which incorporate environmental and social components not measured by GDP.

Disruptive models of degrowth or prosperity without growth are also required to challenge embedded economic path dependencies. Examples include steady state economics, which aims for a balance between production and population growth without exceeding boundaries of planetary health,14 and doughnut economics, which defines the environmental and social space in which inclusive and sustainable economic development can happen.15

Peace and freedom from violence and conflict

Wars, regional conflicts, terrorism, and attacks on human rights activists pose serious threats to peace. For people living in conflict zones, the covid-19 pandemic reduced already constrained mobility and opportunities to establish a new life elsewhere. In 2021 the UN Refugee Agency16 noted that borders were less likely to be open to refugees and that 168 countries fully or partially closed their borders at the peak of covid-19; around 100 of these countries made no exceptions for people seeking asylum.

The pandemic also provided cover for clawing back civil liberties and democratic systems. For example, concerns were raised about the implications of militarised approaches to public health mandates in many countries.17 The global civic society alliance CIVICUS reported that in 2020, 87% of the world’s population lived in countries rated as having “closed,” “repressed,” or “obstructed” civic space—an increase of over 4% in 2019.18 It also identified restrictive legislation to silence critical voices, censorship and restrictions on access to the internet, and attacks on journalists over pandemic reporting in at least 32 countries.37

In 79 countries, security forces used excessive force during protests related to covid-19.

A more peaceful world is requisite to all the other SDGs being met. In turn, if all people’s basic needs were met, rights to democratic expression were respected, inequities were reduced, and we lived within our ecological limits, peace would be more likely. Peaceful relations within and between countries rely on effective, accountable, and inclusive governance systems that can legitimately resolve conflicts between the rights of individuals and those of the community. These systems should facilitate the free flow of information, promote informed decision making and trust in governance institutions during global crises, and provide protected spaces for civil society participation.

Partnership for public good with genuine power sharing

SDG 17 considers “multistakeholder partnerships and voluntary commitments”38 important for achieving the SDGs. Rather than sparking greater multilateral partnerships for health, covid-19 highlighted persisting power imbalances between high income countries and lower income countries.

Covax, a global health partnership set up to provide equitable access to covid-19 vaccines, was initially hailed as an exemplar of global solidarity but was almost immediately undermined by bilateral advance market commitments between rich countries and vaccine manufacturers that prevented it from securing sufficient supplies.40 This resulted in rich countries having much higher covid-19 vaccine coverage than low and middle income countries. The EU, Germany, Switzerland, Canada, and the UK have so far failed to support efforts at the World Trade Organization to agree a temporary intellectual property waiver to reduce barriers to covid-19 technologies, including vaccines.41

The growing influence of the financial sector on global health is also a concern. For example, the covid-19 pandemic saw an increase in the trend to use investment bonds (a form of loan to governments or other agencies) to finance healthcare.42

Investors are also encouraged to support SDG aligned businesses through profit-making socially responsible investments.26 These approaches will inevitably increase wealth disparities since only the wealthy have the capital for profitable investment. WHO’s ability to provide global leadership for health has also been reduced by a steady erosion of its autonomy and capacity over the past two decades,43 and this will likely worsen as it, alongside other UN agencies, seeks private financing to cover inadequate public funding.

The interdependence of all SDGs means more holistic approaches are required, and these will rest on effective public interest partnerships. First, the power imbalances of multi-stakeholder partnerships must be addressed. Private sector interests should not dominate international, regional, national, or local partnerships to realise the SDGs. For example, multi-stakeholder partnerships enable food corporations to participate in the formulation of nutrition policy despite their conflict of interest.44 Public interest, civil society’s voice, and elected officials should instead be prioritised in policy making so that public good is at the forefront.

In addition, effective intersectoral action such as WHO’s “health in all policies” is needed to break down barriers between different sectors and to encourage a joint focus on shared outcomes. Countries will need to establish mechanisms to build citizen trust and adopt collaborative budget processes such as Brazil’s participatory budgeting, whereby residents allocate a portion of public funds to local priorities and needs, including health.45 The public sector must also value, reward, and encourage effective public participation and genuine power sharing.46

Conclusion

Transformative political, social, and economic reforms are needed to disrupt the status quo and promote wellbeing for all. These reforms will enable a redistribution of wealth and power through a fairer economic system based on tax justice and prosperity rather than prioritising growth. They will also reduce carbon dependence to halt global warming and protect and restore the earth’s ecological systems.

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Why is access to respiratory protective equipment still an issue for NHS staff?

The evidence on airborne transmission of SARS-CoV-2 is overwhelming

The World Health Organization and the US Centers for Disease Control and Prevention (CDC) both recognise the airborne transmission of SARS-CoV-2, at short and long range, yet there continue to be delays in implementing respiratory protective equipment across the NHS for staff caring for infectious covid-19 patients. Why?

The evidence on airborne transmission of SARS-CoV-2 is overwhelming, readily available, and expertly communicated. SARS-CoV-2 is an aerosol-borne virus which infects hosts by being inhaled, and this has long been acknowledged in UK government messaging on ventilation.

In the first wave of covid-19 in 2020, the lack of appropriate personal protective equipment (PPE) for healthcare staff was blamed on supply and production issues. This excuse could have had some credibility if one was to ignore decades of pandemic planning that ought to have pre-empted such an occurrence.

However, two years later the official line is that there are no supply issues. So why are staff still not being provided with adequate PPE? It has never been clear if the stated insufficiency of supply is because this has been calculated based on guidance for non-airborne precautions (except for aerosol generating procedures) which, if altered to more sensibly cover physiologically produced aerosols such as when breathing and talking, would significantly alter the levels of supplies required. Given the formal reassurances, this is not a viable reason, particularly in a resource rich country.

Fit tests

Another concern is about fit testing all staff for PPE. While it would have been difficult in the initial stages to fit test all staff, that cannot any longer be the case given that we are two years into the pandemic. Trusts that have switched to using FFP3 masks have, in some instances, done so midway through a pandemic wave (for example, Cambridge University Hospitals NHS Foundation Trust in December 2020), demonstrating feasibility even when the healthcare system was under pressure.

Many trusts have demonstrated that where there's a will, there's a way

Entire staff groups were able to be fit tested over the Christmas holiday period with staff at higher risk of being exposed to covid-19 prioritised and tested within a week. Testing clinics can be opened up before shifts start to enable latecomers or those returning from isolation to be tested and return to work.

Practical considerations include the methodology used for fit testing, either traditional smell testing or the use of aerosol measuring devices (such as Portacount). The former can be challenging as some staff retain the smell of the chemical after a single fail and require a repeat appointment on an alternative day, which makes rapid deployment challenging. Portacounts or similar devices enable both the tester and worker to cycle through multiple disposable options quickly and give real time feedback on how secure the fit is.

Many trusts have demonstrated that where there’s a will, there’s a way, and that it is feasible to implement these processes.

Since January 2021, FreshAir NHS (a group of frontline NHS workers and supporters calling for recognition of airborne transmission and the urgent need for ventilation and PPE) have been requesting that the UK infection, prevention and control (IPC) guidance is updated to accept that airborne transmission of SARS-CoV-2 occurs in all settings and conditions, necessitating airborne mitigations. Initially supported by a petition of 1500 signatories, the request was shared with the UK prime minister and the first ministers, health ministers, and chief medical officers of all four nations. To date we have not received a response even to acknowledge the letter and accompanying rationale for ventilation and FFP3 (or equivalent) provision in healthcare settings. These components are critical, but are not sufficient by themselves as a prevention strategy (vaccination and administrative means are also important); however, any approach to prevention that omits airborne mitigation is woefully inadequate.

Technical debate?

Two years in, there has been very little acknowledgment from the government that there remains an issue with PPE, likely due to the negative press surrounding shortages at the beginning of the first wave; remember nurses in bin liners? On 4 January there was some recognition from Chris Whitty, the chief medical officer for England, when he stated that wider adoption of FFP3 respirators within healthcare “was a technical discussion with several different views on it.” However, denouncing the importance of this problem as a mere technical debate fails to do it justice when so many NHS healthcare workers have died from covid-19 or have longstanding ill health resulting from work acquired covid; or when the bereaved are yet to have answers on why and how this was allowed to happen.

We anticipate that unless airborne transmission is acknowledged, clearly explained, and acted on in infection and prevention control guidance, the discipline of infection control is in danger of relegating itself to obscurity as a credible specialty. High reliability theory, in particular “crew resource management,” dictates when in a crisis, defer to experts and expertise as opposed to simply those in management positions or other positions of power. The NHS needs to embrace the breadth of scientific disciplines and evidence base needed to save lives and minimise suffering caused by infectious pathogens to the benefit of staff and patients.

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Tom Lawton, consultant critical care, Bradford
Matt Butler, consultant physician, Cambridge
Huw Waters, materials scientist
Eilir Hughes, general practitioner

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LETTERS

Selected from rapid responses on bmj.com

LETTER OF THE WEEK

People should be called what they choose to be called

Russell’s letter calling for an end to the anglicisation of our colleagues’ names caused me to reflect on my experience (Letters, 26 March).

I was born in the UK to first generation immigrant doctors from Sri Lanka. For most of my life, my first name has been shortened to the first three letters to make it easier for others to say. This is a throwback to primary school when my teacher announced she could not say my name and abbreviated it for me. I cannot honestly say whether I thought much about whether I liked it or not, but it was clear that it was more important to make things easier for others. I did briefly regret the choice of “Vas” once I was at medical school because of the inevitable jokes.

A year ago, following publicity around the Black Lives Matter movement, I decided to reclaim my full name. I realised that in shortening my name I had anglicised it and in doing so had suppressed a part of my identity. My friends were supportive and took it on themselves to practise the pronunciation. I was touched by their genuine efforts though they did not always get it right. It was more challenging to get colleagues to adapt to calling me something different.

Ultimately, we should call people what they want to be called. In retrospect, I wish I had thought to reclaim my full name earlier and would encourage anyone considering this to do so. I do not agree with Shakespeare when he wrote, “What’s in a name? That which we call a rose by any other name would smell as sweet.” By reclaiming my full name, it certainly felt easier to show up as my true self rather than presenting an edited persona to the world.

Vasandhara Thoroughgood, GP, Colchester

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SELF-CARE INTERVENTIONS FOR HOMELESS PEOPLE

We can reach homeless people in emergency departments

Hopkins and Narasimhan point out that the universal right to health will never be achieved if we continue to underserve the homeless population (Analysis, 26 March).

We need to help healthcare workers provide non-stigmatising care. Medical schools lack training on homeless people and their unique health challenges. Almost all junior doctors have an emergency department rotation, and many homeless people use emergency care as their main contact with medical professionals. This is an opportunity for intervention.

There is a generally poor understanding of government directives around homelessness—if the “severe weather escalation plan” is activated, for example, people can stay in hospital until suitable accommodation is found. Poor communication between government and healthcare workers is entrenched.

Education of healthcare professionals is key to helping the homeless population. Medical contact, most often through emergency departments, provides an opportunity to signpost people towards important resources. We need to do better.

Isabella V C Watts, clinical and research fellow in oncology, London

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LOW BACK PAIN

Gait analysis can help improve lives of people aged 60 and over

I am in the “people aged over 60 years” group described by Traeger and colleagues (Practice Pointer, 26 March). As a result of an incident 14 years ago I had a fracture of the shaft of my left femur that required surgical intervention.

I subsequently developed intermittent right sided low back pain. Physiotherapy followed on several occasions, and recently, when the pain became continuous, I saw a chiropractor for spinal stretching exercises. Those treatments had only a limited, short term effect.

A gait analysis revealed my left leg to be shorter than the right. A heel raise was prescribed, which has led to much improved posture. I can stand up straighter, have achieved the best relief of low back pain so far, and need no analgesics.

Gait analysis was not mentioned by Traeger and colleagues, but it is important to make physicians treating this age group aware of this resource.

Ronald J Atkinson, retired consultant medical oncologist, Hillsborough

Cite this as: BMJ 2022;377:o1059

INTERRUPTING DOCTORS’ WORK

Hospital ward rounds are irritating and ineffective

As a GP with reasonably recent exposure to the ward round, I find it amazing that hospital teams work in this irritating and ineffective way (David Oliver, 19 March).

A group of people going from bed to bed literally looking down on patients. No privacy. Anyone can wander past—even if they don’t interrupt, their presence does. It’s like GPs having their surgeries in the waiting room.

Why can’t hospital clinicians sit in prepared rooms where patients can be brought? They could have a quiet environment, a static computer with access to all records, and the opportunity to develop a meaningful relationship. Bloods could be taken there and then. This might not be appropriate for very sick patients, but then GPs still see sick patients in their consulting rooms. It would be hugely empowering for clinicians and patients, and it might stop ward processes and the public environment hijacking good care.

Graeme Mackenzie, GP, London

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Kenneth Frohnsdorff
Consultant physician in geriatric medicine
East Kent Hospitals (b 1933; qualified King’s College Medical School, London, 1960; MRCP), died from cancer on 27 February 2022
Kenneth Frohnsdorff spent his national service in the Royal Army Medical Corps, attached to the Gordon Highlanders in Aberdeen. He then undertook his junior doctor training at Freedom Fields Hospital in Plymouth, Odstock Hospital in Salisbury, and Jersey General Hospital in St Helier, Channel Islands, where he enjoyed presiding as duty doctor for the Jersey Battle of Flowers. When appointed as consultant physician in East Kent he helped develop the department of geriatric medicine at St Mary’s Hospital, Etchinghill, and encouraged the development of multidisciplinary teams in the stroke unit at the Royal Victoria Hospital, Dover. In retirement he enjoyed his hobbies of archaeology and history, and holidaying with his family. He leaves his wife, Sarah; and two children, Max and Alice.
Sarah Frohnsdorff

Cite this as: BMJ 2022;376:o772

Geoffrey Joseph Bourke
Professor of public health medicine and epidemiology (b 1929; q National University of Ireland, Dublin, 1954; DPH, MA, MD, DHIH, FFPHM, FRCP), died from prostate cancer on 25 September 2021
Geoffrey Joseph Bourke was a senior faculty member at University College Dublin and was known to generations of doctors during their undergraduate careers, culminating in the leadership role of dean of the Faculty of Medicine. However, it was his broader interest in establishing the Faculty of Public Health Medicine at the Royal College of Physicians of Ireland that contributed to professional education and training of public health specialists. He was founder dean of that faculty, which remains the training specialty, but was happiest when sailing his boat. He leaves his wife, Joy, two sons, six grandchildren, and six great grandchildren.
Andrew Seymour

Cite this as: BMJ 2022;376:o768

William Irvine Fraser
Professor (learning disability) Cardiff
University and honorary consultant psychiatrist
Cardiff Community Healthcare Trust (b 1940; q Glasgow 1963; CBE, MD, FRCPsych, FMedSci), died from dementia and old age on 15 February 2022
William Irvine Fraser (“Bill”) achieved a precocious consultant appointment at the age of 29 in Fife. Consultant posts at the Royal Edinburgh and Gogarburn hospitals and editorship of the Journal of Intellectual Disability Research followed. He accepted a chair in 1988 and moved to Cardiff. He was a specialist adviser to the Welsh chief medical officer, assistant editor of the British Journal of Psychiatry, and medical adviser to Mencap. He combined a dry wit with inspirational leadership skills, recruiting many talented doctors into a “Cinderella” specialty, but was happiest when sailing his boat. He leaves his wife, Joy, two sons; and four grandchildren.
Peter McGuffin

Cite this as: BMJ 2022;376:o767

John Samuel Staffurth
Consultant physician
Guy’s Hospital (b 1920; q St Thomas’ Hospital, London, 1942; MD, FRCP), died from old age on 10 September 2021
John Samuel Staffurth (“Sam”) was appointed consultant physician at Lewisham Hospital, with five sessions funded by the Medical Research Council. He set up a simple clinical radioisotope laboratory and experimented with isotopes of sodium and potassium. After several years, Sam transferred fully to the NHS and inaugurated teaching, initially for overseas candidates and ultimately for undergraduate students from Guy’s. He published over 30 papers, predominantly on thyroid disease. At the end of his career he moved to Guy’s Hospital and he retired in 1985. Sam met his future wife, Jean (née Baker), at an interview for a locum post. They married in 1959, moved to Bromley, and Jean became a radiotherapist. Sam died with Jean by his side. He leaves three sons and nine grandchildren.
John Nicholas Staffurth

Cite this as: BMJ 2022;377:o847

Edward Young
Consultant anaesthetist
Reading (b 1936; q Guy’s Hospital, London, 1961; FRCA), died from dementia on 7 March 2022
Edward Young (“Eddie”) was appointed consultant anaesthetist to the Royal Berkshire Hospital, Reading, in 1972 and worked there until his retirement in 1996. Even as senior registrar, he was a superb clinical and scientific teacher with huge concern for trainees’ education and wellbeing. As consultant he became involved with the establishment of intensive care and the implementation of the new operating department assistant scheme. Eddie was a keen runner and continued running marathons until his 60s. He won the Observer’s Azed crossword cup 11 times. He was a gifted raconteur with a host of stories, which often included the eccentricities of his surgical colleagues, all delivered with kindness. Predeceased by his wife, Sally, in 2013, he leaves four children.
Tim Smith

Cite this as: BMJ 2022;376:o773

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OBITUARIES

Cecily Kelleher
Cite this as: BMJ 2022;376:o766
Jennifer McCaughan

Nephrologist and director of Northern Ireland’s Histocompatibility and Immunogenetics Laboratory

Jennifer McCaughan (b 1982; q Edinburgh 2006; MRCP, PhD, FRCPath), died from metastatic leiomyosarcoma on 7 December 2021

Told by a teacher at her primary school that an activity would have to be postponed until the next day, 4 year old Jennifer said, “Procrastination is not good.” Friends of McCaughan believe she never, ever procrastinated.

“She was an outstanding doctor, with a razor sharp mind, and good at passing difficult exams first go,” said Peter Maxwell, former training director for renal medicine at Queen’s University Belfast. “She was at the start of a career that promised great achievements when her cancer was diagnosed in 2020.”

Appointed to the joint post of consultant nephrologist and director of Northern Ireland’s Histocompatibility and Immunogenetics Laboratory in September 2019, McCaughan set about adapting services to cope with the pandemic and improve transplantation numbers. In 2020 a total of 158 kidney transplants were performed in Northern Ireland—more than in any previous year.

As demand for covid-19 testing soared, she rolled out a new way of working at the laboratory, dividing the staff into teams of two, covering day and night shifts and bank holidays. She also introduced training videos. Elaine Boyle, manager at the laboratory, said she and McCaughan were proud of how team members had adapted to different work patterns to undertake covid testing while supporting Northern Ireland’s busiest time for kidney transplantation. “This has been the greatest achievement of the team to date,” she said.

In a tweet the British Transplantation Society described McCaughan as “a committed and energetic colleague, who was a great advocate for her patients and the transplant programme in Belfast.”

Early life and career

McCaughan was born in Ballymoney, County Antrim, the eldest of three children of Jim, a presbyterian minister, and his wife, Alison, who worked in adult education. She went to Hazlett primary school in Articlave, County Londonderry, and then Coleraine Grammar School, where she developed a love of science. She graduated from Edinburgh University medical school in 2006 and received the Scottish Association for Medical Education prize for the most distinguished woman graduate of the year.

She started clinical training in Northern Ireland and in 2010 was appointed to an academic clinical fellowship and started her higher specialist training in renal medicine. “Jennifer was attracted to nephrology because as a junior doctor she had seen the transformative effect of transplants on patients with end stage kidney failure on dialysis,” said Maxwell.

As a research student at Queen’s University Belfast, McCaughan focused on long term complications related to kidney transplants and during her doctoral studies became interested in histocompatibility and immunogenetics (H and I). After completing her doctorate in 2015 she spent a year in Toronto, undertaking a combined nephrology and H and I laboratory fellowship.

“She also had a particular interest in patients deemed high risk for transplant rejection. She provided detailed clinical and scientific advice on how to transplant these patients,” said Maxwell.

In a blog post for The BMJ in 2017 (blogs.bmj.com/bmj/2017/01/26/jennifer-mccaughan-and-ailising-e-courtney-is-balancing-risk-the-most-important-skill-in-clinical-medicine/), submitted when she was a final year nephrology trainee, she urged doctors to be less conservative in assessing the risk of kidney transplants for some patients.

“Objectively, the risk of transplanting highly sensitised patients after a short time on dialysis with a kidney associated with increased immunological risk has a favourable risk-benefit ratio. Subjectively, physicians fear the potential for rejection and graft loss so patients wait. And wait,” she wrote.

Optimal patient management required much improved risk-benefit analysis, she argued. “The capacity to do this effectively for each patient is the cornerstone of personalised medicine and an attribute of the exceptional clinician,” she concluded.

She was a co-author of more than 30 papers on kidney failure and transplantation and was respected for befriending staff at all levels in the organisations where she worked. A person of deep Christian faith, she continued to support renal unit and laboratory colleagues even after her terminal diagnosis.

“She was a short but intensely lived life,” said Maxwell.

Cite this as: BMJ 2022;376:o488