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# Institutional implications for science and industrial capacity: policy lessons from the UK's pandemic response

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Global shortages of critical equipment and supplies induced by COVID-19 forced countries to rapidly build and ramp up their indigenous testing and production capacities. However, the many ways in which institutional and organizational change occurred has not been sufficiently captured. Building domestic capacity requires the leveraging and repurposing of existing domestic scientific and technological capabilities, coupled with intensified global outreach to new and existing partners and suppliers. Using the framework of institutional variety, this paper looks at two facets of the UK's COVID emergency industrial response: (1) building its laboratory testing capabilities and (2) for increasing production of personal protective equipment; assessing the institutional capacities and relations that were leveraged in this regard. It uses these findings together with observations of 'innovation processes under emergency conditions' and the potential uses of a 'critical equipment policy' to sharpen some of the recommendations made in the UK's post-COVID Research and Development Roadmap.

**Keywords:** industrial policy; institutional variety; capacity building; COVID-19; UK; science infrastructure.

## 1. Introduction

The COVID-19 pandemic has underscored the importance of national autonomy in responding to emergency conditions and the freedom to make complex choices about medical equipment and testing, with local health outcomes in the foreground. However, this has been done in a context of highly uneven, conflicting, global health guidelines, and inter-country information sharing about how to respond to COVID-19. Rather, with little industrial policy expertise, the WHO issued a universal guideline to 'test, test, test' with understandable interest in, but incomplete and uncertain, clinical foreground detail and very little attention to the vital industrial background and national customization needs of countries as they moved to initiate testing in a supply-constrained environment (Srinivas, Prasad, and Rao 2020). Policy needs to acknowledge that in different national and regional contexts, the co-evolution of key institutions regarding industrialization and healthcare is not one of uniformity but of institutional variety (Srinivas 2021, 2023). Understanding this variety of how institutions (customs, norms, guidelines, standards, laws) and organizational forms can change during an emergency can be critical both to crafting and implementing an effective pandemic response and aligning emergency health response with long-term economic development gains in the development of core innovation processes and critical equipment.

Equally crucial was the political economy of building such capabilities. While shades of a new economic nationalism

pervade much of the political discourse in support of the domestic capacity building, such rhetoric distorts both the real policy and industrial response to COVID in most countries. Our findings point to a different process in which institutional variety (IV) is useful to explain how and why an inward focus on domestic capacity and production cannot be separated from, indeed is coupled with, intensified global outreach to new and existing suppliers worldwide. Contrary to some of the accompanying rhetoric, the actual policy and practice is one where no country could do it alone. In this perspective, the pandemic illuminates the adaptability through recombination of industrial innovation systems and the continued importance of external sources of knowledge and resources, with new emergent strategies under emergency conditions. While this industrial policy response under highly compressed timelines could be viewed as largely temporary, the details of the recombination and extent of IV matter to its post-pandemic UK industrial strategy and future emergency responses in a variety of national contexts.

As such, this paper looks at the COVID emergency industrial response of the UK, specifically its Lighthouse Lab Network, and its efforts at increasing the production and supply of personal protective equipment (PPE). Both laboratory-based diagnostic capability and PPE production demonstrate innovative repurposing capabilities under rapid change, and constitutes a crucial example of a critical equipment policy that could have long-term economic development potential if recognized as such. In the case of increasing testing capacity, we show that the UK leveraged and expanded both its existing

domestic laboratory capacity and university–industry relationships, with the help of incumbent international partners and newly established global suppliers. For increasing access to PPE, the UK pursued a strategy that aimed to increase domestic production of PPE through non-traditional suppliers of PPE in the UK and through aggressive contracting with both existing and new global suppliers of PPE. In this sense, the UK epitomizes this dual track industrial strategy, where the immediacy of the pandemic forced the UK to rapidly leverage domestic scientific and business engineering capabilities while seeking secure links to global suppliers for the same types of capabilities and products.

For this paper and its IV perspective, we take a preliminary look at the institutional relationships and resulting capabilities that shaped the UK's COVID emergency response; this informing potential future research on the nature, governance, and speed of such change, a crucial question for the organization of scientific efforts and industrial policy design. Finally, we look to put forward a set of broad policy recommendations that both inform and reinforce the UK's post-COVID Research and Development Roadmap. The UK is a compelling case to begin our inquiry. While the UK's COVID response was national, strategies for building domestic capabilities and securing access to PPE were, on the one hand, coordinated between the four nations of England, Scotland, Wales, and Northern Ireland. On the other hand, each of the devolved administrations exercised varying degrees of autonomy in implementing the national strategy.

This paper is structured as follows. [Section 2](#) builds our research framework and key assumptions. [Section 3](#) explains our methodology and the case study approach. [Section 4](#) and [5](#) present our two cases, along with some initial analysis regarding key assumptions. Findings from each are further discussed ([Section 6](#)), including implications for policy and future research. [Section 7](#) concludes.

## 2. IV, innovation, and emergency response

National responses to COVID-19 have generated interest among academics and policy makers as to how the institutions that govern and produce innovations have adapted and changed in meeting this necessary response, particularly in the push to develop new therapies and vaccines. For example, [Sampat and Shadlen \(2021\)](#), taking an innovation systems approach, argue that the US response to COVID-19 has involved a change in government funding of biomedical research, from a pre-pandemic focus on basic research funding and patent-based development incentives to a new emphasis on late-stage product development and procurement agreements. Given less attention in the literature, although not less important, are expected changes to other areas of the biomedical innovation and industrial production system as a result of COVID-19, notably in the areas of diagnostics and other medical equipment. Whereas new therapies and vaccines will initially require significant R&D, an emergency ramp-up for diagnostics and other essential medical equipment will likely involve less intensive R&D and more emphasis on rapid development and deployment of capabilities and scale-up processes. Our choice of framework and attention to PPEs emerge from an interest in industrial systems and the R&D functions and manufacturing embedded within it.

### 2.1 IV, emergency response, and organizational interactions

Building upon innovation systems, yet diverging in fundamental ways, IV is an approach that considers how different institutional contexts and relationships over time shape the way new knowledge is sourced and converted into productive capabilities (see [Srinivas 2021](#)). ‘Institutions’ means the evolving set of customs, norms, guidelines, standards, and various rules, that shape and govern innovation and related industrial policies. For example, industrial procurement, intellectual property, good manufacturing practice standards, are all examples of institutions in industry, but so too are new, more fluid, hybrid public–private partnerships, special purpose vehicles for investment, consortium bidding, and so forth. Specific organizations and agents act under such institutional frameworks, and also shape institutions in turn over time, a feature well recognized in scholarship of institutional as well as evolutionary economics. In this way, IV can demonstrate and explain how the institutions that structure, facilitate, and govern innovation and new organizations—behave rather differently in different national, regional, and sectoral contexts, and in this paper, under emergency stressors.

Moreover, the health policy response itself can be framed through the institutional emphasis on specific industrial institutions. For example, [Srinivas, Prasad, and Rao \(2020\)](#) highlight at least seven biological and clinical uncertainties that depended on exactly how countries industrially approached the problem of Covid-19 diagnostic kits production, making the relationship between laboratory science and clinical effectiveness highly dependent on available industrial characteristics and the organization and network ability to operate at scale, a point that WHO health policy directives did not appear to recognize. Furthermore, the several uncertainties on biological and clinical ends and their industrial dimensions are not directly equitable and are also parsed differently by professional communities well before PPE or diagnostic kits supply meets demand. In essence, countries with existing technological capabilities when faced with trade blockages and/or lock-downs had to scramble to institute customized scientific as well as engineering problem-solving, and to institute systems of incentives and norms and laws that ensured diagnostic kits and PPEs were rapidly manufactured and used to clinical specifications.

Similarly, in looking at emergency response to communicable disease outbreaks (e.g. TB, Malaria, and Ebola), [Rama-lingam \(2015\)](#) posits that under emergency conditions, the innovation process can look much different than it is normally described. First, because of the rapid unfolding of most emergencies, timeframes for making decisions are condensed. Second, available resources are also limited, and this difficulty can be compounded by the location of the emergency. Third, processes for innovation and processes for delivery will need to occur concurrently. Finally, these accumulated pressures often result in solutions that rely on known technology and processes rather than on new technology and new approaches to either production or delivery. Overall, the need to rapidly respond to the said emergency constrains the innovation process, forcing it to make decisions on a narrow, more familiar set of options and processes that are more likely to address the emergency in an acceptable timeframe. As such, the focus of the response is on the scaling-up of existing capabilities and the repurposing of know-how, rather than the development





had long-established links to global suppliers for reagents and testing equipment for which they relied on exclusively, i.e. there were no domestic UK suppliers. That said, the UK did not have a working test and trace system in place, although was attempting to develop such a system. As UK and global cases rose significantly in February and March, the UK's 40 NHS labs were carrying out about 5,000 tests a day, far short of what was needed: the UK simply did not have the lab capacity for large-scale public testing. Furthermore, no reliable testing system had been developed or implemented and, most concerning, global supplies of needed reagents and swabs were limited due to the unprecedented global demand (Baraniuk 2020; Kirkpatrick and Bradley 2020). As a result, the UK had to do the following: (1) increase laboratory capacity; (2) create a testing system (3) secure future testing supplies; (3) implement an effective test and trace system; and (4) do this rapidly (1–2 months). All four challenges were pursued. However, to do this, the UK placed most of its efforts toward expanding and developing laboratory testing capacity and re-establishing global supply links to source critical testing supplies and equipment (UK DOHSC 2020).

#### 4.1 Building laboratory testing capacity: the Lighthouse Lab network

For building its laboratory testing capacity, the UK initiated plans to increase the capacity of existing NHS labs (NHS and Public Health England)—from 5,000 to 25,000 tests per day and to create 3 Mega-labs (Lighthouse labs) designed to boost mass testing to 100,000+ per day, along with tests at drive-through centres and at homes. For its choice of sites, the UK took advantage of a range of life science investments over the last decades (UK DOHSC 2020). The three lighthouse mega labs are:

(1) Lab in Milton Keynes (opened on 9 April 2020) at the offices of the UK Biocentre, a not-for-profit business established in 2011, established and funded by the UK National Institute of Health Research;

(2) Alderley Park in Cheshire (opened on 20 April 2020), run by Medicines Discovery Catapult Ltd, and funded by Innovate UK. It is located on what was once the laboratories of Imperial Chemical Industries (ICI) (see below) and what is now a science park;

(3) University of Glasgow lab (opened 24 April) located in its Clinical Innovation Zone at the city's Queen Elizabeth University Hospital campus, funded as a Scottish Catapult by the Scottish government (UK GOV 2020a).

The Lighthouse Lab Network (LLN) also includes the establishment of three smaller labs for regional and complementary capacities: a lab in Northern Ireland run by Randox, a Cambridge-based lab run by AstraZeneca and GSK, and a lab in Newport, Wales, brought online by PerkinElmer. Both AstraZeneca and GSK provide support and expertise to the entire network, with supplies and equipment provided by ThermoFisher Scientific, TECAN and Brooks Laboratories (UK DOHSC 2020). The labs are supported and governed by NHS England, Public Health England, and the UK Department of Health & Social Care, with governance of the Glasgow lab led by the Scottish government (LLN 2020).

Looking at the LLN overall, there is significant diversity among the various labs in terms of each site's origins and

pre-COVID use, the extent of industry collaboration, and local actor involvement (the latter to be discussed later in the paper). The lab at Alderley Park is a good example. A historically important site for pharma research in the UK, Alderley Park was the site of the pharmaceuticals R&D laboratory of the UK national champion ICI. ICI was originally set up after the First World War, an early state investment towards a private company to help catch-up to German chemical giants. This became Astra-Zeneca during the Thatcher industrial 'reforms' which led to ICI's division. When Astra-Zeneca closed Alderley Edge and moved to Cambridge, it became a science park for spin-off and new companies (Pharmaceutical-technology.com 2021).

The origins of the Milton Keynes lab are of more recent history in that it is built as an expansion to the laboratory facilities of the UK Biocentre, a non-profit organization established in 2014 (UK Biocentre 2021a). As another example, the Glasgow lab is located in the newly developed University of Glasgow's Clinical Innovation Zone, close to the Queen Elizabeth University Hospital. In this way, the Lighthouse labs, for the most part, are all built upon and/or are expansions of existing pharma and innovation-based sites, aligning with aspects of our assumptions (2) and (3), positing that the time urgency of emergency situations will force nations and localities to build on existing capabilities rather than develop inherently new capabilities.

Despite the organization diversity mentioned earlier, there are a number of additional institutional commonalities among the labs. First, the UK government and devolved administrations play the lead role, as expected in assumption (1), both in terms of funding and coordination, with Public Health England, NHS England, Department of Health and Social Care, and their Scottish equivalents all taking the respective leads. Second, the three mega-labs are managed by governmental or non-commercial entities. For example, the labs at Alderley Park and Glasgow are governed by Innovate UK and government-funded Catapults. Furthermore, the Glasgow and Cambridge labs in particular, have strong university linkages and support, including the use of university faculty, support staff, lab space, and equipment. Also, the three main labs are all housed either in public or non-commercial sites and facilities. Additionally, all main LLN sites became operational within 4 months of the pandemic's emergence in the UK. Again, government and public involvement in these sites is significant, this includes government and publicly owned facilities and capabilities, lending support to assumption (1) pertaining to the essential and leading role that government plays: it is government that has the mandate and resources to so rapidly ramp-up testing capabilities in this way.

While led by government, the LLN can be described as a partnership between government and private industry, with industry taking the lead in both providing the bulk of laboratory equipment, particularly in new equipment provided for expansion purposes, and for overseeing and managing, to varying extents, the day to day operations of each site. For example, AstraZeneca and GSK not only play the lead role at the Cambridge lab but are also significant participants in the other LLN labs. Both long-established pharma giants have a strong presence in the UK, with GSK having its global headquarters in London. Providing laboratory instrumentation and reagents to the LLN is US-based Thermo Fisher







- FFP3 masks: Alpha Solway re-shored mask manufacturing from Taiwan and increased production; Don & Low imported and installed new machinery to manufacture filter material for masks.
- Eyewear (visors & goggles): Alpha Solway switched emphasis from making protective clothing for oil and gas industries to visors. Also producing eyewear were: 4C Engineering (an off-shore engineering company based in Inverness) and Aseptium (a decontamination company) & Lifescan (a J&J company); Skyrora (a Scottish company involved in rocketry), and Baker Hughes (a US oil drilling equipment company).
- Aprons: Berry BPI, part of US owned Berry Group, is a major UK-based plastics and rubber company, and Europe's biggest plastics recycler, already a supplier to the NHS, sourced and shipped to Scotland specialist machines for the manufacture of disposable aprons from their Greenock factory.
- Non-sterile gowns: Don & Low (a Scottish company owned by Thrace Group (Greece) repurposed production to produce material for gowns. These materials were then converted to gowns by Edmund Bell (Yorkshire) and Keela (Glenrothes), with additional support from Endura and Transcal (Livingston).
- Ventilators: JFD Ltd Aberdeen and Inchinnan leveraged their expertise in breathing equipment to design a new ventilator; Babcock's Zephyr Plus ventilator is being supported by Plexus and Raytheon.
- Hand Sanitizer: CalaChem Ltd produced sanitiser at its site in Grangemouth, with ethanol provided by Whyte & Mackay: whisky distillers.

In looking at both the UK and Scottish CEP, our findings very much support our assumptions regarding innovation under emergency response conditions, particularly in terms of institutional leadership, government and industry partnering, and the leveraging of regional capabilities. First, for the UK's CEP and that of the devolved administrations, government has taken the lead role in both implementing policy and coordinating the procurement and production strategy (assumption 1). For example, Government leadership seems critical in selecting and courting manufacturers who are not traditional producers of PPE, i.e. without government support and championing, the uncertainty of such a change of operations would be too much for most companies to take on. Second, the UK and Scottish CEP are clearly two-tracked in that they seek out sources of PPE external to the UK while leveraging national and local PPE capabilities (assumption 2). What is interesting about the UK's CEP efforts, and is different than its testing capability strategy, is that it is less about leveraging existing PPE production capability and more about developing new PPE production capability, this through the 'Make' strategy. Finally, the fact that devolved administrations implement their own CEP and leverage, as the Scottish case exemplifies, their own, often regionally based capabilities, lends some support to our third assumption that *institutional interactions toward developing capabilities will occur most prominently at the local and regional level and that it will be local or regional capabilities that will be leveraged in this regard (assumption 3)*. This opens up questions or future research of what national industrial policy can or should do to boost the cohesion of existing, dynamic industry efforts.

## 6. Discussion: future research and policy recommendations

### 6.1 Findings and open questions

Overall, our findings regarding the UK's response to both increasing capabilities for COVID-19 testing and diagnostics and production and supply of PPE lend considerable support for our main assumptions about the importance of detailed analysis of local conditions to understand the institutional context for innovation under emergency conditions. That said, there are aspects of our assumptions that were not easily captured by our research approach. First, while our findings demonstrate a COVID-19 strategy where 'emergency' conditions require both rapid decisions and development ramp-up of capabilities, we do not capture an assumed 'speeding up' of learning processes, either institutional or organizational.

In other words, questions regarding how public and private capabilities were so quickly integrated are left unanswered and may require more attention to institutional change within the State (e.g. centralized discretion and a small group of decision-makers which might otherwise have been wider, slower, consultation). Second, our findings do not lend much insight as to whether and how local and global capabilities interface. We assume that much of this takes place at the regional level, but we do not find overwhelming evidence for this.

Future research could address both areas—learning processes and institutional coordination—in more detail, especially around which capabilities and locations are policy-induced to cope with the assumed ability of scarcity-induced innovations. Therefore, a fertile area for future research is on the learning processes, outcomes, and implications for the institutions and actors involved in both the LLN and the PPE 'Make' strategy. For example, what have firms such as Scotland's BioAscent and India's Brooks Laboratories, gained and learned by participating in the LLN and what might this mean for future emergency response efforts when crucial inputs are scarce? Also, not captured in our research findings is evidence of changes to regulatory processes, i.e. if regulatory approvals of new equipment were sped up (e.g. PPE), how were required safety and technical standards met; how have such process changes influenced post-COVID-19 regulatory practices? In some ways, answering these questions will take a more micro-oriented approach, one that looks more closely at organization and programme-specific decision-making and learning processes that should shape future industrial policy design, i.e. starting at the micro level and then connect to and build up to a more accurate institutional view.

### 6.2 A new variant of import substitution industrialization?

Earlier in the paper (Section 2.2), we suggested that national responses to COVID-19 through building domestic capabilities and selective external outreach have some similarities with traditional import substitution industrialization (ISI) as well as 'catch-up'. However, recognizing that only some institutional combinations are possible in emergencies (which could be induced by any stakeholder but most likely by emergency policy responses), and only some of these combinations are possible because of the presence of existing technological capabilities, we can provide some nuance to the concepts when tested in emergencies. Also worth noting is that ISI is not occurring in the traditional sense i.e. crucial imports





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