# OPUS2 

Scottish Covid-19 Inquiry

Day 10

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(10.00 am)
        (Proceedings delayed)
(10.05 am)
THE CHAIR: Good morning, everybody.
            Right, now, unusual today because we're going to
    have the first witness remotely and for that purpose we
    have to have somebody who will do the vetting of it, so
    in case there's any mistakes, as those of you who were
    here yesterday will remember, we can deal with it that
    way, and because the only way we can get a camera that
    can see whatever it requires to see for the YouTube
    channel, Mr Caskie is sitting up here beside me.
MR CASKIE: I've not been promoted!
THE CHAIR:So that's fine. With that introduction,
        Mr Caskie.
            Witness HSC0037 (called)
            Questions by MR CASKIE
MR CASKIE: Can I firstly check that the witness is able to
    hear me?
A. Yes, good morning.
Q. Good morning. It's nice to see you. I should say at
    the beginning there is a restriction order made by
    Lord Brailsford to ensure that basically no one is named
    in the hearing today. So the hearing is about your mum
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and your dad and I'll simply be referring to them as
    your mum and dad and there is also reference to a care
    home and I'll be referring to that simply as "the care
    home"; okay?
A. Okay.
Q. If you could do likewise, that would be very helpful.
        As I understand it, your mum was born on
    9 September 1927 and she's now -- is she now 96 or is
    she about to become 96?
A. She was }96\mathrm{ in September just past, yes.
Q. Your father was born on 25 April 1928 and he sadly died
    on 30 May 2022; is that correct?
A. That's right, yes.
Q. Okay, firstly, can you tell us about your mum before she
    became ill?
A. Mum was, I would say, a very traditional mum. She was
    a teacher. She taught science at secondary school and
    she loved nature --
Q. I'm sorry, can I interrupt a second? The volume in
    here --
THE CHAIR: It's my fault. Can you all hear? Is there some
        equivocation about that? Yes, someone at the back
        shaking her head. I wonder if we can get the volume
        turned up. Could you ask the technical people if we
        can --
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MR CASKIE: Well, the technical people are listening so they will be running through even as we speak.
THE CHAIR: Sorry, I'm giving any plenty of indications that I'm not very good at technology. It's justified.

Are they going to see what they do? They're fixing. Right. I apologise for that delay, but we'll see if it's better now.

You were asking about the witness' mother.
MR CASKIE: Yes, I was asking you to tell us something about your mum. You said she was a science teacher.
A. She was a teacher and worked, as well as bringing up her or their three children, myself and my siblings. She loved nature, she loved gardening, she knew an awful lot about the scientific names of plants. As well as working, she was quite a home bird, I would say. She loved cooking and baking and dress-making and knitting. I don't think I've picked up many of those skills from her. She was quite sociable, she loved sailing, when she was younger, in dinghies. She loved dancing, she loved playing bridge. She was very sociable and a lovely mum.
Q. Can you tell us a bit about your dad?
A. Dad worked for the same company all his life, which I think is quite unusual these days. He was a research scientist. Dad was also very sociable. He loved

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spending time with his family and spending time with his friends and blethering perhaps over a whiskey or two. He had a great love of literature, he had a great love of history and he was particularly interested in Scottish history, Scottish music. He spoke a little bit of Gàidhlig and was very proud of his Scottish heritage.
Q. Now, I understand you live in South-West Scotland?
A. I do, yes.
Q. And latterly your parents lived in North-West Scotland?
A. They lived there for about 30 years, yes.
Q. On Google Maps it tells me that you lived about 220 miles apart.
A. Yes, that might well be right. It was certainly about five hours' drive one way.
Q. And I think towards the end of your father's life both your mother and your father were in the same care home.
A. Correct. Mum went in first and Dad, for a while, sort of managed at home with the help of carers and then, latterly, he was in the same care home as Mum, and that's where he passed away.
Q. Okay. Now, l'll ask you a bit more about that in a moment, but, as I understand it from the witness statement, your mum entered the care home in the spring of 2018.
A. That would be right. She'd been in the local hospital
Q. You said that your mum had gone into the hospital for about two months.
A. Yes.
Q. Who was looking after your dad during that period?
A. I moved up for a while to be with him. I was very worried about both Mum and Dad and so for a while I spent half my time with my father and then half my time back where I lived. While I was there, I did as much as I could to look after Dad, as well as taking him much as I could to look after Dad, as well as taking him
to visit Mum, and so I sort of commuted between the two places.
Q. It's a long commute.
A. It is a long commute.
Q. Tell me about your dad's health problems.
A. Dad was $--I$ used to call them Tweedledee and Tweedledum because Mum had severe physical issues and Dad had increasingly severe mental health issues with vascular dementia. So, latterly, when they were at home together, Mum would do a lot of the planning and thinking and Dad would do as much of the doing as he could. They sort of managed very well. As I say, they
for about two months with ongoing infections and during
that time her physical health deteriorated to the extent
that she couldn't -- she wasn't able to return home to Dad.
Q. Was there a specific incident that precipitated her going into hospital?
A. Yes, yes. They were just about managing to be independent at home. They were fiercely independent. They didn't take help easily from social care. They'd never -- we'd never had any interaction with social care until the very late stages of my parents' life. And Mum, who had osteoarthritis, had real difficulty walking, must have tried to get out of bed one evening during the night to the commode that was in her room downstairs and she took a fall and, unfortunately, wasn't found until the carers came in that morning and she was on the floor of the room. And then she was taken to the hospital and actually after that she wasn't able to return to her home or to Dad.
Q. You said that your mum went into the care home in the spring of 2018. Is it correct that your dad went into the care home in 2019?
A. Yes. So Dad for a while managed somehow, with some struggles, to remain at home. As I'd said earlier, they were both very home-orientated and he managed one way or
the other for a wee while, while Mum was in the care
home. Unfortunately, with his dementia, he --
Q. Well, I' II come back to that.
Q. You said that your mum had gone into the hospital for
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were very independent. I'm sorry, I've forgotten the question. I'm sorry.
Q. You've answered it. It's fine.

So you've got a mum who has physical problems and a dad who has dementia problems. Between the two of them, they made a complete person?
A. They did. They managed so well between the two of them. They did have some support from the home care team that would come in and, certainly for Dad at that stage, they would administer Dad's medication -- some of the home care team were responsible for administering his medication, which was very, very important, and would also look after some of his needs. He latterly wouldn't make food or drink for himself so they would try to make sure that Dad had meals put in front of him. He didn't always want to eat them, but they did their best.
Q. I think you said that your mum was in the hospital for two months after her fall.
A. She was there over two months. She just -- she had sepsis -- she had ongoing infections, sepsis, urinary infections, and they just couldn't seem to clear it for her.
Q. What happens at the point at which she's getting out of the hospital in terms of her care?
A. We didn't actually know what to do. Where they lived,

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there wasn't a huge number of care homes or respite beds available, but it was very obvious that Mum at that stage really couldn't stand, couldn't walk and was starting to have even restricted movement when she was sitting down or in bed. And I think it was at that stage that I first had my first contact with the social care team, who were exploring her going into a care home and were trying to find a care home that was in the vicinity of where my father was so that obviously they could continue to see each other.
Q. Was that the primary selection category for the care home that your mum was going into?
A. I'm sorry, I don't know --
Q. That she was close to where your father was living?
A. At one point it seemed that wouldn't be possible. They were keen to move her out of the hospital and there wasn't, as I said, an immediate availability close to where my father was. But then, as these things tend to happen, a bed became available in the care home that was local to where my father was and so very quickly that all swung into place. She was very reluctant to go but really there was very little option for her.
Q. Did your father have wider family living in the vicinity?
A. Yes, yes, he had.
Q. Without naming them, can you tell me what relatives they were?
A. My father had many uncles and so he had cousins, first and second and a few third cousins very close and on hand, so he was in familiar surroundings where he lived, and, yes, there was quite an extended family on hand for him.
Q. So we know that your mum goes into the care home and your dad is still living in the family home. He has cousins in the vicinity. Are the cousins involved in some way in the care of your father?
A. Yes, yes. The days when I wasn't able to be with my father, helping look after him, I actually had a WhatsApp group of some of the core family, I guess you'd say, and I'd ask them to help by maybe taking Dad to see Mum or maybe doing some particular shopping for him or taking him for one of the many medical appointments that he needed. So we depended a lot on local family and a few close friends who were also in the area.
Q. In the area, was he well known?
A. Yes, very well known and well respected and well liked.
Q. Over time, did his condition deteriorate?
A. Yes, it did. It did, and I was so worried about him. When I arrived to see him, if I was going up to stay for

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three or four nights -- and I'd go in the back door and he'd be sitting in the room by himself, sometimes sleeping, and, you know, Mum's chair would be empty. So he managed very well and he never complained, he never really complained, but he was always glad to see somebody when they walked in the door. The family would just knock and walk in the door and he was always glad to see people when they came in. But I really did worry about him being in the house by himself because at that point he was worried about Mum, he thought she was still in hospital, he never quite grasped that she was in the care home, so he did start to deteriorate.

The carers would leave a sandwich for him at lunchtime and they would return and the sandwich would be uneaten or they would make a cup of coffee for him and return and find the coffee was cold. He increasingly -- he always used to get out of bed early, but increasingly they'd arrive at maybe half nine in the morning and he'd still be in bed, which was not like him. He was always an early riser. So even at that point, Mum went into the care home, I think he missed her dreadfully and was very lonely because they had been so used to looking out for each other.
Q. Did anything in particular happen to his weight?
A. Yes. He really wasn't eating very much at all and we
were concerned about the weight loss and he would always
brush it off. As I say, he was never one to complain and he would say, "I don't have much of an appetite. I'm not doing very much these days. I don't go out and about. I don't have much of an appetite", but he was eating very little. The carers phoned me one day to say he'd nearly fallen down the stairs because his trousers were so loose on him that even with a belt they'd nearly fallen down and he'd nearly tripped down the stairs. They were quite steep stairs and they were just very concerned at that point that he was losing weight. He was getting more frail and, although he had never complained, I think the carers were starting to feel that he was getting weaker and more frail and really wasn't able to look after himself.
Q. When those events were developing, did social work and the medics basically make a suggestion as to what should happen with your dad next?
A. Yes. I think there was a difficult week or so when I - because of work, I think, I really couldn't get up there for about ten days and I had been going up pretty much every week. It was all a bit crazy, but there we are. We do these things. He was starting not to even wash his hands, wash his face. He'd stopped shaving. We couldn't get him to a hairdressers and he started to

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look quite weak and dishevelled and there was a suggestion that he went into respite care, which -- as I understand it, he was entitled to two weeks' respite care. It was suggested that for his own well-being he would go into a care home for -- initially for two weeks' respite care, which was such a relief to know he would be looked after.
Q. At that stage in someone's life, it 's quite often the case that the social work department or the doctors will seek to have the person assessed in terms of their mental and physical capability and, quite often, people go into hospital for a few weeks to do that. Did that happen with your dad?
A. Yes. Yes, it did. That was actually before he went in for respite care. As I say, he had vascular dementia and he was starting to display -- he'd had it for many, many years. It was a very slow -- a very slow, gradual decline. He was quite canny about dealing with it and, to be honest, hiding it as much as he felt he could, but because his behaviour started to change and he was uncharacteristically cross and -- he started actually hallucinating and so he was taken into a geriatric psychiatric residential $--I$ find that hard to say -geriatric psychiatric residential --
Q. Unit, yes.
A. -- hospital for assessment, which was in a different part of Scotland. So again I was then trying to drive to see Mum and then drive to a different location to see Dad. I was hugely concerned at that point, but with hindsight that was a good thing because he then had a full psychiatric assessment of his dementia, which he'd never had before. Really at that stage he had had very little help with his dementia because they were seen to be managing one way or another. So, yes, he went into a geriatric hospital for I think about ten days.
Q. At the conclusion of that ten-day period, was the suggestion that he simply return home?
A. It was. It was. He was given some medication for the dementia or for the impact of the dementia, which actually did seem to pacify him and keep him calm, so he returned a bit more to the lovely, lovable, kind-hearted character that he had, and he returned home for a short while, but, as I've said before, he really then continued to deteriorate, so -- but by then he'd had the full diagnosis and I think it was on his record that he was becoming less and less able to manage in his home by himself, even with my care and even with the home carers going in. He was almost getting beyond that.
Q. So at that stage was it decided that he also should go

## into a care home?

A. I think at that point then they suggested that he go in for respite care. It was such a relief. It actually meant for a wee while I didn't feel I had to go up all the time because I knew he was actually in 24 -hour care and, amazingly, he was in the same care home as Mum, and that was just -- he actually thought he was in a hotel, but never mind. He was in the same place, the same building, as Mum and that was just lovely. They would sit together. Sometimes they would both be asleep in the armchairs but at least they were together. It was such a relief to the family to know that they were back together and, to be very honest, it was a relief for me to know that I really didn't have to go up all the time and arrange things for Dad and try and look after him.
Q. So both of your parents are now in the same care home.
A. $H m m-h m m$.
Q. Can you describe the care -- well, sorry, your dad went in for two weeks' respite. Did he ever come out?
A. No.
Q. No?
A. No.
Q. So the respite became his permanent residence?
A. Yes. Again, it was just the way that care homes work, that beds become available and it was just the way that
a bed became -- a permanent bed became available for Dad. Some people said, "Are they in a double room?", and I said, "Care homes don't work like that. They don't have double rooms". They were in individual rooms but they could spend their day beside each other, which was wonderful for the family and for them.
Q. Was it a nursing home or a care home?
A. I believe it was a care home, so that there were carers rather than nursing staff, I believe.
Q. And can you describe the home, just physically?
A. It was a small care home. It was a care home that was local to where my parents lived and where my father's family were. I felt very fortunate, once I'd started, as many of us did, to learn more and more about care homes. It was a small care home. There were some local staff there who also had some knowledge of local history and the local area, which -- Dad thoroughly enjoyed talking to them. He really blossomed in the care home for a short while because he had the social interaction that he so much enjoyed. I was always happy with the care that Mum and Dad were given, always happy. There were some very dedicated carers that went above and beyond and, until COVID struck, I was so happy with the care and it was such a relief to know that they were both together again and some of the care, I -- you know,

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was given to them and, as I say, the family and -- we didn't have to organise so much for them.
Q. Tell me about the staffing levels in the care home at that time.
A. At that time it was so difficult for the care staff. There were often shortages of staff and the staff would often do extra hours to fill in the gaps out of sheer goodwill and dedication. I was very grateful for that. They would often call in bank staff, who were sort of on-call, as it were, to come in, so that would be people that would come in to cover shifts. I believe the situation with the shortage of staff was because there was a shortage of local housing, so staff -- people couldn't move into the area to take up the jobs because there was a shortage of housing, particularly affordable housing or rented housing or local authority housing, and that was across the whole health and social care in that part of North-West Scotland, that, because of the shortage of affordable housing, it meant the whole healthcare system really was struggling with a shortage of staff. And that definitely affected the care home and at times I think they struggled with cover and it was only the goodwill of the staff to do extra hours that they always, one way or the other, managed to get -- the great thing was I could go in and I could
help -- sometimes help Mum to eat or I could sit with Dad and keep him company or I could do a jigsaw with Dad or I could help Mum. So at that time, you know, I was happy to go in and help look after Mum and Dad the way I always have done. So -- but there was a shortage of care staff, even before COVID.
Q. At paragraphs 15 and 16 of your witness statement you talk and name a number of staff at the care home. I'm going to ask you not to name any of the staff. But reading those two paragraphs short, it appeared to me that the management of the care home was in a state of flux. Would that be a fair summary?
A. It was, it was. It was like a double whammy that, just as COVID was starting, there were, for different reasons and different circumstances, two or three managers across the period of pre-COVID and into COVID, again just through circumstances, and that was -- that didn't help the situation during COVID, that there were different people coming in and -- yes, there were different staff, different managers, coming in.
Q. When your dad went into the care home, what were his immediate needs? I mean, he had just come from hospital.
A. Dad had left the psychiatric hospital and was at home for a couple of weeks before he then went into respite

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care. As one carer said, he looked a bit dishevelled, he looked a bit dishevelled. He needed a good shave, he needed a hair-cut, he needed a good feed, and they looked after him, they tempted him with different food, and he sort of made friends with one of the other elderly gentleman and they would sort of sit --
Q. Did you see an improvement? Did you see an improvement after he arrived?
A. He did, he did. A short-term improvement, but I think he so much liked being where Mum was. He loved the social interaction. Latterly, when he was at home, he didn't really get out and about that much. He became quite insular and withdrawn, and now he had people around him that he could talk to. He enjoyed word games and dominoes. Even with his advanced dementia, he still retained the capacity to play dominoes and he enjoyed blethering with some of the other residents who had a similar connection with the area that he did.
Q. You described yourself travelling up to see him frequently. What about your siblings and his wider family in the area, would they go to visit your parents?
A. Yes, yes. So some of his extended family would go in and take him gifts. I don't know what gifts, shortbread, the odd ... my siblings live even further away than I do so I tended to be the sibling who took
A. Almost overnight. The doors were locked and that was that.
Q. And people that he did see, presumably they would be wearing masks?
A. Yes. All the carers $--I$ mean, it was awful for the carers being in what was quite a warm building wearing masks and to this day I think that Dad's last year

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really, he rarely saw a smiling face. With his dementia, he never quite understood why people were wearing masks and he would always -- almost every day he would say "Why are you wearing masks? Why are you wearing masks?", so he never understood.
Q. Were your parents hard of hearing? Had they become hard of hearing?
A. Dad's sight and hearing were quite good. My mum, even before she'd gone into the care home, was slightly deaf and had combined macular degeneration and cataracts. So even before the care -- before she went into the care home, she had hearing and sight difficulties and the masks just made that ten times worse.
Q. Well, tell me about that. Why were masks such a significant interference?
A. They really limited Mum's communication or Mum's hearing. Even with her eyesight -- she I think learnt some element of lip-read or she could look at people's expressions and she was quite sharp mentally, Mum, until latterly, so she picked up on people's expressions, even though her eyesight was limited. But the hearing -- we had to really raise our voice and speak quite loudly to get her to hear, and with the mask, you know, that became more difficult.
Q. Those visual cues as to what someone was saying, they
A. By the carers wearing a mask.
Q. Aha, and also later visitors ?
A. Yes. Yes, she was continually having to say, "What?", "Pardon?", "What are you saying?", so it just became quite a struggle to say anything. We persevered but the masks - - I think, for my dad, not seeing a smiley face -- and, as I've said, Dad was such a sociable person - - and not to see faces and for my mother not to be able to readily hear conversation and hear what was being said to her made everything very difficult.
Q. A moment ago we referred to carers and also visitors. How did you regard yourself, a carer or a visitor?
A. I wasn't a visitor, I was a carer. I was a care-giver in the true sense of the word. I for years had looked after Mum and Dad as best I could, living away, but I would spend a lot of time looking after them, organising their care, giving them their care, helping them with all sorts of personal care, emotional care, well-being care. My brother had power of attorney but he made it very clear to the social and healthcare -the health team that $I$, in his place, was his deputy, so I considered myself a care-giver. I cared for them and I looked after them and helped them and was happy to do

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## so because they looked after me.

Q. So at this stage both parents are together in the care home but visits have stopped?
A. Yes.
Q. Were any effort made to try and organise technology to allow you to speak, FaceTime or anything like that?
A. A lot of people were FaceTiming at that time, I think, not just those in care homes, but friends and family all over the country were FaceTiming. That was very alien to Mum and Dad. They didn't have broadband in their house. They didn't have mobile phones. They just weren't the generation that used technology. So everybody was saying to me, "It must be great that you can FaceTime Mum and Dad", but I couldn't. The care home's wifi was very weak and very intermittent and really only worked in the main office area, where I suppose the modem was, and Mum and Dad couldn't go into that area because it was a communal area. So although we tried initially, it just wasn't possible. The wifi signal wasn't strong.
Q. And who was responsible for the wifi?
A. I never quite understood. There seemed to be some to-ing and fro-ing between the local authority and the local NHS because of the status of the care home, and they seemed to -- when I asked about it -- pass
responsibility to one or the other and then I was told by email that the care home wasn't a priority to improve the wifi, given the COVID situation. There were other - - a local authority, maybe the hospital or -that the wifi was more of a priority. It was very difficult. We tried to get wifi set up, but Mum would fall asleep while it was being set up, Dad could not understand what I was doing on the television, I couldn't hear, they couldn't hear, and then the wifi would collapse and that was that.
THE CHAIR: Can I ask a question about that? I'll be very careful because I don't want to disclose anything I shouldn't. But you've said that your parents' care home was in the north-west of Scotland. Was it in a geographical area where wifi was, as a matter of generality, difficult to obtain or there were obstacles to it? I say this as someone who lives in a rural area so I know about these problems.
A. Yes, yes. Where the care home was and where my parents had lived would exactly be in that rural area where wifi was limited. The availability of wifi providers was limited and the strength of the signal was limited.
THE CHAIR: Right. So might I say that there were possibly -- and I understand fully that you don't have the detail of this - - but possibly wifi connection

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presented more of a problem in the area where the care home was situated than it would have, for example, in Edinburgh or Glasgow?
A. Initially, yes. Yes. As the pandemic went on, it was realised that there were ways around that, with boosters and other technology.
THE CHAIR: Yes.
A. And it eventually improved, but, initially, yes, the wifi signal --I eventually asked for us not to do the FaceTime because Dad would get upset, Mum would get upset, I'd get upset, we all got upset, and it was just worse than not trying. So eventually I said to the care home, "I can't - - I can't put Mum and Dad through this", so initially it caused more problems and more upset than any of us wanted.
MR CASKIE: In February 2021 I understand that changed to some extent. I'm looking at paragraph 25 in your witness statement.
A. It took a while, it took a while, but, yes, that's right, February 2021, this wonderful thing called "MiFi" was introduced. I'm really not sure where it came from.
I think it was through some charity. And that made
a difference because then it was so easy to get on to
wifi and, amazingly, having said that Mum and Dad
weren't really into technology, Dad very readily
accepted that I was on the television screen and Mum readily accepted that I was there, and it was great because I could see them. I could see that they had a piece of cake or they had some shortbread. I could see what jumper Mum was wearing. Mum could see me. And it made such a huge difference that I could actually see them and talk to them, even though we still had to sort of speak quite loudly. It was an amazing difference.
Q. Did you regard the year that you were waiting, as it turned out, for the wifi to be installed as a wasted year?
A. I lost a year of communication with Mum and Dad, yes, and I worried about them. They must have worried about me because, as I said, I was the one that kept them up to date with all the family news and they weren't getting that, so they were not only geographically isolated, they were isolated from news about their family. Phone calls were difficult because Mum couldn't hold the phone because of her osteoarthritis. She couldn't hold a phone and the carers would prop it up on her pillow and it would fall away and -- so a whole year they lost news and contact and connection with loved ones and we lost connection with them.
Q. I'm still working through the statement, but I want to try to take the next part of your evidence relatively
shortly. You indicate that you were keeping track of all guidance issued by the Scottish Government.
A. Yes.
Q. How? How were you doing that, initially and subsequently?
A. I was - I think, like many of us who were separated from loved ones, I was looking a lot at websites, particularly the Scottish Government website or websites. I found that hugely confusing. I like to think I'm fairly au fait with technology and business-speak and terminology, but I found it very confusing trying to find the specific piece of guidance or the specific piece of information that referred to the care homes, and it was at that point that I stumbled upon or I was referred to or somebody told me about Care Home Relatives Scotland, and to this day I --
Q. Did you find that a useful resource?
A. Night and day. Hugely. Hugely. And I've said time and time again, I feel for the many hundreds, thousands, of people that never found Care Home Relatives Scotland. They were such a fantastic resource for information. They clarified the guidance, they provided support, and time and time again I found information clarified through them that I would have struggled to from the Government website, which I found very confusing.

Endless links and just ... so Care Home Relatives Scotland, night and day, what a difference it made to me.
Q. Now, we know that there was a process -- and I'll ask you about the specific process in a moment -- but a process by which visits were introduced. We said earlier that you lived about 220 miles from where your parents were living. Was any allowance made for that 500 - mile round trip in terms of visits once they were authorised?
A. I don't think it was, to be honest. The -- as I say, the care staff were always wonderful, but the restrictions that were placed on them meant we had to book an appointment to see Mum and Dad, and I had to phone and make an appointment and make sure that I was there for the appointment. I did ask for weekend visits. I did --I'm self-employed so I was doing some work from home during lockdown and I had some flexibility . But when I was busy with work, I wanted to travel up at the weekend or maybe drive up on the Friday and go home on the Monday and visit Mum and Dad at the weekend. But I was told specifically for that care home that wasn't possible and it had to be during the week, almost like a 9.00 to 5.00 visit, and that was difficult for me.

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Q. Why was that? Were you given an explanation as to why that was?
A. I did query it and I had and still have a good relationship with the care home. I was told it was because they didn't have the full staff at the weekend, so maybe some of the catering staff or the cleaning staff or -- the staff weren't there, so the carers that were there had to take on extra duties and they couldn't therefore be there to let me -- to unlock the door and let me in and sit with me or do my COVID test, and so I had to try and manage my time to be there during 9.00 to 5.00 during the week.
Q. Now, at paragraph 30, you describe a visit.
A. Yes.
Q. Can you tell us about that visit? Firstly, was that a formally arranged visit in terms of the guidance that had been issued?
A. Yes. I think this was when -- this was the first time I actually got to see Mum and Dad in some sort of face-to-face capacity. When I say "face to face", I think this was the time of the window visits and Mum was on the ground floor and she actually had a small single door to the outside, and so I would again have to book the visit and several times -- and I've got photos -- I would be standing outside the door, about
a metre away, Mum and Dad were about a metre inside the door, Mum was swathed in blankets, Dad was covered in blankets. Being in that part of Scotland, it was always blowing a hoolie and often raining. I would be in my cagoule. I had to wear a mask.

It was a small opening, I was about 2 metres from Mum and Dad, I could see them, but with the mask and probably the wind and rain, I was having to sort of raise my voice. I was glad to see them but it wasn't a satisfactory visit because I couldn't reach out to them. Dad kept saying, "Will you not come in?", Dad kept saying "Come and have a cup of tea", Mum kept saying, "You look cold, come on inside", and that was all they wanted to say, was "Come on inside". And although I could see them, I couldn't really have a conversation with them because I had my mask and we were 2 metres apart. They were inside, I was outside. I suppose I was reassured that I could see them. Then Mum would say she was very cold and she'd have to go back inside and then Dad wanted to go and be with Mum and I'd have to go away again.
Q. I get the impression from what you're saying that they were unsatisfactory, the visits, at that stage, although there was a great relief to see them.
A. I did those visits because it was better than nothing,

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but in a very strange way almost the Zoom calls that we did were almost better because I could communicate more easily and it was a more warm atmosphere for Mum and Dad. So I did continue those window visits but I didn't feel we could communicate easily. I didn't feel we could have conversations. It wasn't satisfactory, no.
Q. At the time, did you think what you've just said, that the FaceTime contact was better than seeing them in the flesh in terms of your communication?
A. Well, the window visits or that open-door visit -I think they were sometimes called "garden visits". It wasn't a garden. It was just like the back of the care home - - yes, in a way it was easier because Mum and Dad were more relaxed. Mum could even be in her bed and Dad would be sitting beside the bed, so Mum was more relaxed, Dad was more at ease. We could have a conversation. It was quieter. In a strange sort of way - - although I was so grateful to see Mum and Dad, in a way I could communicate better with them over the Zoom calls, the FaceTime calls. However, I did continue to do those garden visits for a while because it was --
Q. Did you continue to do the FaceTime as well?
A. Yes, yes, we still did FaceTime. So I would maybe visit them during the week and then have a FaceTime call at the weekend, so I did my best to try and keep up the
contact because I was worried that they would think we'd all forgotten them.
Q. You say that the next thing which happened was being able to sit outside with them.
A. Yes. So at some point Dad would come outside and I would sit, again, a wee bit apart from him with a mask on, but I think this was - - the weather had improved by this point and I could sit near Dad outside. It was difficult for Mum with her osteoarthritis. She was in great pain, she didn't like being in the wheelchair, it was uncomfortable for her and she could really only tolerate about 15 minutes in the wheelchair. So, again, although it sounded great that they could come outside and sit with me, Mum could really only sit for about 15 minutes and then she would go inside. But at least I could sit with Dad for a bit and blether away to him, which was lovely. It was lovely.
Q. The next thing, the next development, was visits in the conservatory.
A. Yes.
Q. Tell me about those. Tell me about the physical layout of the place the visits took place.
A. You know, there $--I$ feel for the care home at these different stages because they really had to work the best they could to organise things. There was

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a conservatory that Mum and Dad could enter through the care home and I would enter through the outdoor. There was a perspex plastic screen between us and I would sit on one side of the screen with a mask on and Mum and Dad would sit on the other side of the screen, Mum was in the wheelchair, and they would have a cup of coffee and some cake or biscuits with them. Again, it was good to see them, but Mum again could only tolerate about 15 minutes in the wheelchair and Dad just didn't like being behind the screen. He kept asking the carers to move it .
Q. You also say that you had a concern about there only being one visitor allowed, ie yourself.
A. Yes, there was $--I$ had a strong understanding that the guidance said "Up to two visitors", and sometimes my husband would drive up with me, and that was great because he could do the drive and I could just sit there and it took some of the strain away from the drive for me, but he wasn't allowed in at the same time as me and that was very difficult because he and Dad got on so well. And Dad would ask - - I'm sure Dad probably thought we'd got divorced or something because he never saw my husband and he would ask after him and -- you know, it was just -- Dad just loved to see people and during that time that was another relationship that he
had gone -- he was very fond of my husband and he asked after him and Mum asked after him and yet he was never there, so it was difficult. And I had understood the guidance said that my husband could be there but the care home said that that was not possible, so that didn't happen.
Q. So you were receiving conflicting information?
A. I think the guidance was so confusing and -- yes, so it was conflicting. I never blamed the care home because they were doing their best to try and interpret it, but the guidance was very confusing. There was no clear instruction about what to do. There was endless links and you had to go to different pages and this had been updated and that had been updated. I certainly didn't want to cause any friction with the care home. I didn't want to cause any problems because they had Mum and Dad's life, so I just accepted it.
Q. You say -- I'm looking now at paragraph $38--$ that you requested essential visits in relation to your father on 19 February 2021. What was the response to that?
A. Do you know, I don't actually -- it was through Care Home Relatives Scotland that I'd heard about essential visits. I honestly don't recall any communication from the care home about that, that wording. I was concerned that Dad was getting more

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vague. He was more withdrawn. He would give shorter answers to my questions and he wouldn't follow up with questions of his own. So he'd stopped asking about the wider family, he stopped asking about what I'd been up to, he stopped referring to other members of the family. He was just getting more withdrawn and he looked more vague. He looked like he was becoming more -- yes, more vague and I just felt he was -- his mental awareness was declining. He just wasn't himself, he was getting more withdrawn, and at that point I asked if I could come in and make a visit and I don't recall a reply to that.
Q. You say something about infection control. At paragraph 43, you make reference to different approaches being taken to infection control.
A. $\mathrm{Hmm}-\mathrm{hmm}$.
Q. Can you tell me about your experience of that?
A. This was to do with the testing, the COVID testing. Again, I was very happy that Mum and Dad were being tested and the staff were being tested, whatever the regime was then. I of course would always test before I left home because I wasn't going to make a five-hour drive and then find I had a negative test, so I would always test before I left home and then the care home would ask me to do a test once I got on the premises. Different staff would handle it differently. So some
staff would want to administer the test, my test, themselves, so they would put the swabby thing either up my nose or into my mouth. They had masks on and gloves and aprons; some of the carers would sit and watch me do the test ; some of the carers would ask me to sit in the car park and do the test; other carers would let me sit in that conservatory area and do the test; some of the carers would just say, "Have you done the test?", and I'd say "Yes"; some of the carers would ask to see evidence from the COVID app thing to say that you'd submitted the test; some of them didn't. It was quite a wide variety of different ways of checking whether I'd done the test.
Q. So there was quite a lot of inconsistency?
A. Yes. Yes. I mean, I myself was always very certain that I had tested and got a negative test both before I left home and at my visit, so I was very happy, but there did seem to be a range of different methods used by the care home staff as to what they were doing. I was happy to go along with it because I knew at the end of that, whatever method they were doing, I would get in to see Mum and Dad, so I just went along with it. But, yes, different care home staff had different ways of testing.
Q. Now, during this period where you're having some visits,

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are any siblings getting any visits at all?
A. One of my siblings didn't visit at all. He found the whole thing very distressing and upsetting and didn't visit at all, and I would relay photos and messages from him and he would speak on the phone, but he didn't. My other sibling, the one with power of attorney, did make some visits. He had a considerable travel to make and again he was working and he had a fairly essential job, you might say, during COVID and -- but he made some visits. But I was the one that did most of the visiting .
Q. What do you think the impact of that isolation from his family was on your father and mother?
A. They never outright complained because they were the generation that didn't complain. They were quite stoic about things. But they would worry. I could see -- and they said they were worried. They didn't understand what was going on. As I say, Dad became more withdrawn and vague and then Mum worried about my father. They worried about each other. I think the lack of contact -- particularly for my father who enjoyed social contact, I think it was particularly difficult for him. Even when I wasn't visiting, I made reference I think to the fact that he had some -- he made friends in the care home. He had to sit apart from his - - they weren't
friends before he went in. He made friends with some of the residents. He would have to sit apart from them. They couldn't sit round the same table and have their meals. They would have to sit at separate tables. They couldn't play their dominoes. They had to sit apart from each other. I think that had an awful impact on him, that he didn't have that social contact with anybody during those last stages of his life.
Q. And then you said at the outset that some time later your father died. We have the date of that.
A. He did. That was after the Open with Care came into play, the next endless stage of whatever was happening on the guidance, and he had a very rapid decline. The family, his close family, had stated that we'd rather he wasn't transferred to hospital unless it was an absolute emergency and the way of his decline meant that he died in the care home. He was well looked after. I was grateful to the staff.
Q. At the end of your witness statement you have a section headed "Lessons to be learned" and in paragraph 53 you essentially say emails are not as good as phone calls; is that your view?
A. No, absolutely. Absolutely. We did get fairly frequent emails and I've passed them all to the Inquiry. I struggle to look at them now. There was so many of

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them and often there were emails referring us to the guidance and I would be looking at it and thinking,
"That's not my understanding of the guidance", and the guidance can be interpreted different ways. I would phone at times the care home, sometimes --I was only allowed to speak to the manager. The carers were asked always to refer to a manager rather than -- both my parents had specific carers but we always had to speak to the manager, but, as you'd said, there were changes in management staff. They were so busy at that time. They were short-staffed. They themselves said they had endless video team meetings with Infection Control and Public Health and NHS and Social Health, so it was really hard having a conversation and trying to find -even if I'd said, "I'II phone at 2 o'clock in the afternoon", I would phone and find that there was an emergency meeting or somebody had been called or something had happened. I really felt it was difficult to keep up to date with Mum and Dad's care. I was restricted on how I could visit and an email saying, "Your mum and dad are fine, they're doing well", that kind of email just didn't give me any insight into how they were.
Q. When you did speak to them, did it seem that they had time to talk to you?
A. They didn't have time. They were rushing from one thing to the next. They tried -- the manager tried, but they were rushing from one thing to the next and it wasn't through lack of want but they just did not have time to give me an explanation of how Mum and Dad were doing, and, again, I felt so cut off from my mum and dad's care.
Q. I'm now looking at paragraph 57, where you talk about new guidance coming out and then it taking time to filter down.
A. Yes.
Q. What was your experience of that?
A. I knew the guidance before the care home seemed to hear the guidance, so there were one or two occasions when I would be phoning up and saying, "I believe I can now do a garden visit", "I believe I can now come in" or, wonderfully, "I believe I can actually now come into Mum's room with all my PPE on", and the care home manager would say, "We haven't yet had the latest guidance" or "There's a meeting on Thursday when we will have the nearest guidance". So, again, because of my attention to the Scot Gov website and because of Care Home Relatives, I often knew about the changes before the manager did. And no disrespect to the manager, but she would have to wait until there was a meeting or

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several meetings had taken place before she then had a - - when I say "meeting", I mean a Teams meeting. So I often knew the guidance before they did.

No criticism of the care home, but it took about a week for it to be filtered down and then it would often take some time for the care home to make the arrangements, like setting up the plastic screens or setting up the visitors ' or care-givers' tests. So it took a while. It wasn't always instant.
Q. Could you read paragraph 58, please, aloud?
A. "Throughout the pandemic I was treated as just a visitor."
Q. Before you came in, you told me that if you get upset,

I should push you to finish.
A. Yes.
Q. I'm pushing you to finish.
A. Yes. It's three sentences:
"Throughout the pandemic I was treated as just a visitor. I was my mum and dad's carer and [I was] an important part of their care. We shouldn't have just been removed from our relatives' lives."

I shouldn't have been removed from my mum and dad's life.
Q. Thank you very much. I don't have any other questions for you apart from one. Do you feel as though you've
said everything that you need to say?
A. I do. I feel I've had the chance to care about Mum and Dad in telling this story and I think that's my role as their carer, to tell their story and to tell our family's story, because I am their carer. I was my dad's carer. So thank very much.
MR CASKIE: Thank you very much.
THE CHAIR: Yes, thank you very much indeed. Thank you.
Right. We'll take a break now until 11.25 .
MR CASKIE: Perfect.
THE CHAIR: Very good. Thank you, all.
(11.08 am)

> (A short break)
(11.26 am)

MR CASKIE: Could you bring the witness in? One more witness this morning, sir. In fact one more witness today.
THE CHAIR: The witness' name?
MR CASKIE: The witness' name is Kristin Duncan.
MS KRISTIN DUNCAN (called)
THE CHAIR: Ms Duncan, please take a seat.
A. Good morning.

THE CHAIR: Good morning. When you're ready, Mr Caskie.
MR CASKIE: Thank you.
Questions by MR CASKIE

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MR CASKIE: Could you tell Lord Brailsford your full name, please?
A. I'm Kristin June Duncan.
Q. The Inquiry has details of your address and date of birth and so on. You have very helpfully provided a witness statement for the assistance of the Inquiry extending to I think 24 pages. Before you signed that, had you read over it?
A. Yes.
Q. And are you happy that the content of it is true?
A. Yes, I am.
Q. And although you'll provide some oral evidence today, you would want Lord Brailsford, I assume, to take account of all of the parts in your statement?
A. Yes, I would.
Q. Good. We're here principally to talk about your mother.
A. Yes.
Q. Who was ...?
A. Catherine Christina June Scott.
Q. Known as ...?
A. June Scott.
Q. And she was born in -- I can't read my writing -13 June 1929?
A. 9 June 1929, yes.
Q. 9 June -- sorry about that -- and died on 22 March 2021?
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## A. That's correct.

Q. Okay. We're going to talk about her care and ultimately her demise, but before we do that, tell us a bit about your mum.
A. Well, Mum lived to a grand old age of 91 years and for most of her adult life she was a mother and she reared her four children, but she juggled that with a career in PE teaching. She had qualified from the Irene Marsh College of Education in Liverpool in about 1950 and she taught for most of her adult life, along with balancing the act of bringing up a family. But she was also passionate about Scottish country dance and she will perhaps be remembered as a great teacher and adjudicator of Scottish country dance.
Q. In that, I see from the witness statement at paragraph 7, she led a group of Scottish country dancers at the Jakarta Highland Gathering.
A. Yes. Yes, she presented candidates for the Scottish Country Dance Teaching Certificates, and that was back in 1982, and she was also appointed as an adjudicator to the Royal Scottish Country Dance Society, so it was through that role that she was given wide responsibility for adjudicating and teaching in Scotland and England and her adjudicating took her to very exotic places like Jakarta.

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## THE CHAIR: Highland Dancing, the same thing?

A. No, no, sorry -- a very big difference, Royal Scottish Country Dance Society and Scottish country dancing, yes. THE CHAIR: What's the difference?

## A. How long have you got?

THE CHAIR: Well, we're not going to ...
MR CASKIE: Can we go to paragraph 10 of the witness statement, please? Can you just read that?
A. Yes.
"In 2008 she was awarded the highest honour from the Royal Scottish Country Dance Society - - The Scroll - for her encouragement of others to appreciate and enjoy Scotland's heritage of dance and music."
Q. And that played an important part in her life?
A. A huge part. I mean, she'd been teaching Scottish country dance since earliest years, when she was just a postgraduate, so that's all of her adult life, until of course arthritis took over, and she took this group at Nethy Bridge for 19 years and was very fondly remembered by them for her contribution, but eventually arthritis took its toll.
Q. And she lived with her husband in Badenoch \& Strathspey?
A. Yes.
Q. Tell me about the home that they made there.
A. Well, they retired to Badenoch \& Strathspey in the late
duty", but obviously with great love and affection we gave this duty, and we would set out our rota for the full year so we knew which weekends we were actually doing to help with all the things that she needed to let her stay at home alone.
Q. So that's her weekends taken care of. During the week what was happening in terms of support for your mum?
A. Yes, so quite quickly she had to be reassessed by social services. Both my parents had had -- they were on the radar of the Social Services team and they had been receiving free personal care. There were things they needed help with. So she was reassessed when she was on her own and gradually over the years -- and I'll try to give you a picture that concertinas that -- but over the years gradually the Social Services had to keep on reassessing her and add in more and more help.
Q. So it started out, as I understand it, at two days per week?
A. She was getting a visit two days per week -- no two visits per day, sorry. Two visits per day. When she needed help with things like getting up, getting dressed, taking pills, remembering to eat, prompts, that's when the Social Services put in home carers. So we had the Care at Home team at first and then Highland Home Carers latterly. These two sets of teams,
Q. What happened then?
A. Well, my father died suddenly of a heart attack and -unexpected really, but that left our mother now widowed and a huge challenge then of being on her own, living alone in a very rural setting, about three miles outside Kingussie.
Q. After she was widowed, did she get support to continue to live at home in the community?
A. Yes. I mean, this was her wish and we very much rallied round her from the word "Go". We recognised, with her not being able to drive anymore, her challenges of living alone multiplied, so we, the four of us -- the four children - - we worked through what we needed to do and we provided support for her at weekends essentially, because we worked, but we actually managed to keep a system going for the next seven years and we devised a rota where we set out our -- it was called "Mother
1980s and they created a dream retirement home. It was
rural setting and had large grounds and was really responsible for the construction along with the help of builders and also the garden. He played a huge role in that. They lived there very happily and looked after each other well until their old age.
Q. Until 2012? A. Well, my father died suddenly of a heart attack and -11 13 14 15 17 18 19
20 21 22 23 24
getting to the periphery; hands, feet, and she was at risk of things like amputations. We were always concerned that might happen. It didn't ever. We were lucky. She was lucky. But one of the other things with PVD, as it's called, is vascular dementia. So she was getting that and it was not rapid or aggressive, it wasn't of the Alzheimer's type. It was vascular dementia, which meant that gradually her cognitive skills were declining and also memory loss and ability to do things and balance. So things were going and that's -- it was slow. We would see it gradually increase over that sort of seven-year period.
Q. I'm asking this question as a kind of placeholder -I' II come back to it - -
A. Yes.
Q. --but what was happening to her hearing?
A. She had suffered hearing loss because of probably primarily the PVD for quite some time and had been given hearing aids and had a lot of help from the services that provide hearing aids, but invariably with elderly people, they lose them, they don't like them, they get background noise, it doesn't work, and she stopped herself eventually, didn't want them, couldn't be bothered with them, and with so many other things to 49
deal with, almost the hearing was -- it was in there but it wasn't the top priority.
Q. And what about her -- well, talking of top priorities, what about her ability to walk, to remain vertical, not fall ? What happened to that?
A. Well, she had always been of a strong build. She'd been a PE teacher and she carried herself very well. Deportment was her thing with the country dance. So she was a strong lady and despite that she was getting old and frail, so then falls became an issue and she would fall over small things in the home. We lifted carpets and mats to try and make sure she didn't, but they happened, and carers would find her prone on the ground when they came to do their visits. And of course this became more frequent as it got ...
Q. And in terms of the carers and presumably also the doctors in the vicinity, did they suggest a way forward for your mum ultimately?
A. Yes, ultimately. Ultimately the carers would have a logbook in the kitchen and they would fill in every visit what they saw and what happened so we were able to read and see what was going on. We knew what was going on because very often I was called or my brother was called. She had the red help button. We were going out, you know, increasingly in the middle of the night
to answer things. And so --
Q. Do you live in the vicinity?
A. I was seven miles away and my brother was about 17 miles away.
Q. And you were getting increasingly frequent calls?
A. Increasingly, and others who were on the list because we had a list of six people who would answer that call. So it was increasingly difficult and, to cut a long story short, ultimately we tried to keep her at home for her 90th birthday, that is what she wanted. That was where she wanted to be.
Q. That was her target?
A. That was her target and we could see her -- and we knew from the Christmas that it was getting nigh-impossible, so we teetered on the brink for six months with this.
Q. And during that six-month period, did you do anything about finding somewhere for her to go?
A. Denial in our heads, hoping that miraculously it might not be needed but realising it would be, thinking about it, I would have a look at places locally without actually going to them, but nearer June, when she had her 90th birthday and the writing was on the wall, clearly on the wall, I did go and look at several in the vicinity and did my search further than what I would have liked, but I just wanted to get a feel for what

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they were like.
Q. In the vicinity was important, I get it .
A. Absolutely, absolutely .
Q. Why? Why for your mum was the vicinity important?
A. Well, she had established a fantastic network of friends in Badenoch, she had done an awful lot for Arthritis Care in Badenoch and had -- with her teaching hat on, she would take classes -- even when she could no longer do Scottish country dance teaching, she would take a class for elderly people, frail people - maybe not so elderly, but people who were suffering from lack of mobility. She would teach whether they were sitting in their seats or standing. She would give them mobility and fun.

So she had a huge network of friends, from her country dancing, from Arthritis Care, from people in the village in Kingussie. You know, they were all there. So we felt, if she's going to have to go into care, let 's get somewhere where these people can still visit her, because they would.
Q. And in terms of selection, where did you select?
A. There wasn't a big choice. Badenoch \& Strathspey is a very long, narrow, old county. There are two or three main towns in it and there were only three care homes in the district. So we didn't want to go outwith Badenoch
\& Strathspey, where her heart was, where her home was, where her people were and her friends, so we had a limited choice.
Q. Can I take you to paragraph 24 in your witness statement?
"We put Mum into care against her will however. This was devastating, both for us and us [as] children."

Let me ask you a couple of questions about that. Did she need to go in?
A. Yes, she did. We had no choice. We had a strong steer from the medical practice that it was -- time was up and she was at risk at her own safety of staying at home alone any longer.
Q. You said that she had a target, which was to stay at home until her 90th birthday. Did she get there?
A. She did. We did it and she went into care on the 19 th. Her birthday was the 9th.
Q. So ten days after?
A. Yes.
Q. So she worked to her target?
A. She did. She did. That would be typical.
Q. Again, I'm going back to the statement. Can I take you to the end of paragraph 27? What you say there is, just the final sentence:
"Our parents had been very organised in how they
sorted out what was to happen to everything in the event of their demise/death."

Then you begin that same paragraph by saying:
"With regarding to DNACPRs, both my parents had put that in place many years previously and I believe this was held by the Kingussie Medical Practice."

And then it gets transferred to the care home. So they had made plans for their future?
A. They had, they were very organised, and I remember the day when we were all called to have a meeting with them to talk about this, all these aspects, and of course it was hard to take it so seriously then but we did and they made us take it seriously. And I would have to say they were just highly organised in that respect.
Q. At paragraph 28 you talk about a deterioration in your mum's condition. Tell us about that.
A. Well, yes, she had been going in and out of a day care centre in Kingussie for the last three or four years for day care and therefore, when she was put into somewhere new, completely new, all of a sudden that was a bit confusing. Her memory loss -- sorry, her memory wasn't that great. So, I mean, she wouldn't have been all that aware of time and she wouldn't have remembered who had popped into see her necessarily when she was at home, so going to a care home, it would have added to the
confusion, but there was confusion there anyway because of short-term memory loss, if that makes sense.
Q. At paragraph 29 you say that the next eight months were an adjustment period. Was that really your mum coming to accept that this was where she was going to be?
A. I think she didn't like it. She always said, "When are you taking me home?" or "Take me home". That was always a request or a plea. So did she accept it? I don't think so, but she had to adjust to it .
Q. Now, in terms of visits into the care home, let's start with how much freedom you had to visit.
A. $\mathrm{Hmm}-\mathrm{hmm}$.
Q. How much freedom did you have to visit?
A. It was very good, excellent. I think the care home was very enlightened in its approach to ensuring that the family felt that this home for their loved one was their home as well and we were welcomed with open arms. We had access to the pad -- the pad for the key to the door, so there was no restrictions on that. We could visit when we wanted to, any time of day.
Q. You say, "We could visit when we wanted to". Who are the "We"?
A. We as a family, so this was her four children, but also her grandchildren and her great grandchildren and any friends from the ones I've described previously from her

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life further up the valley, from the Arthritis Care or her neighbours, her nieces, her gardener. Her hairdresser even came down from Kingussie to see her and her home help. Anybody and everybody whose lives she had touched, who were within travel distance -- and even nieces from Edinburgh came. You know, anybody could come and visit her in the care home.
Q. Could you read paragraph 32, please?
A. You want me to read it?
Q. Yes.
A. Yes:
"Mum's health slowly deteriorated, nothing new, just gradual deterioration, she was after all slowly reaching life 's end. Had it not been for a strong heart, or perhaps a zest for life and a will to continue to see her family, all of whom she always recognised, she might perhaps have passed away sooner."
Q. At paragraph 36 you say that you received an email from the managing director of the company that owns the care home. What did that tell you?
A. Paragraph 36?
Q. Yes.
A. So this was -- yes, the pandemic was upon us and we were all very aware of that across the country and the care home actually had already -- my sister tried to visit on

13 March and she was - - she phoned in advance and they said, "No, don't come". So there had been almost like a two-week closure, if you like, of relatives, friends being told not to come. But 20 March was a singular moment when we did receive an email from the managing director saying that there was to be a lockdown from that day, not unexpected, but apart from that there was very other detail -- there was very little detail.
Q. And at paragraph 37 you talk about the care home putting in place attempts to facilitate communication by iPad. Did that work for your mum?
A. No. Absolutely it couldn't. It never would. She was old school. She had never had a mobile phone herself. She didn't know what an iPad was. And she -- they were great actually. There was an activities co-ordinator who really, really tried to get us to have contact with her using an iPad, but she couldn't understand what the talking head was on an iPad, so this didn't work and we gave up on it. We agreed to give up on it very early on.
Q. Tell me about communications from the care home to you and the wider family.
A. So generally, in the first couple of months of the pandemic, when nobody really could move anywhere and we were all confined to our houses -- and I think there was

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a lot of acceptance around that state as we waited for things to change -- the only way to communicate, to find out how she was, was literally to phone up and ask. So we would do that and phone and just say, "How is she? How's she getting on?", so that there was always an ability to make questions and ask about her welfare.
Q. And what responses were you getting when you were phoning up to say, "How is my mum?"?
A. So if the person who happened to answer the phone at the time had just seen her earlier that day, that would be great. They could say, "Oh, yes", and they would give you a quick report. But nine times out of ten you would get somebody perhaps who was maybe doing more of an admin job or who hadn't been out on the wings and they would very helpfully say, "Yes, well, look, I don't know, but I will get somebody to call you back who does know", and they would. They would call back and give you that update.
Q. And that would be every time you made contact with them?
A. Yes. I didn't ever experience not getting any feedback when it was asked for, but also very conscious of there was only so much you could expect them to know or tell you on the spot, coming in cold like that. We did assume that if there was any emergency or any big change, then they would contact us.

# A. I hadn't heard anything in the news and I did watch 

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every day to see what was being said, and so, therefore, I made an assumption that this was just an initiative that the care home took to facilitate a visit outdoors, and, as it transpired, that's exactly what that was because I got that in May. I was invited by phone by the activities co-ordinator, suggesting if I came down, they would wheel Mum out and we'd be able - I would be able to speak over the railings at a distance of 2 metres. And the actual guidance did come out from the Scottish Government on 25 June. That was the formal published guidance. So this was like a good month anyway before there was published guidance, but we got it .
Q. You were on the other side of a fence. Was that the boundary fence?
A. There were several boundary fences to the property. It looked out on to lovely fields with ponies in it, a rural setting. This was kind of a side part of the care home, rather than the main garden, but nonetheless it was an open space, a green open space, and it had railings.
Q. How did your mum react to that meeting?
A. Well, it was very difficult because it was the first meeting I'd had $--I$ hadn't seen her for well over eight weeks or longer, maybe ten, and she hadn't seen
any family member in that period and she was angsting straightaway about "Taking me home". And because she was peering at me through a railing -- because she was sitting and the railings were at least a metre -- I was crouched down to see her at her level, so she was seeing me through a railing and there were cars behind me so she actually thought she was on a railway platform, bizarrely. There are no railway platforms where she lived. She didn't remember that or know that. She just thought she was on a railway platform and that I'd come to take her home. And the whole conversation was around going home and trains, and it was circuitous.
I couldn't get her off this -- and very difficult
because I couldn't get near her, I couldn't give her
a hug. I couldn't reassure her in any way.
Q. She was talking about going home?
A. She was talking about going home, and the trigger was the railings and the cars behind.
Q. At paragraph 43 you say:
"Copious guidance for Care Homes emerged from the Scottish Government ..."

Tell me how you think that worked, the issuing of guidance at that stage in terms of how the care homes dealt with that?
A. Yes, there's a couple of things in that. I think the

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copious guidance actually at that time, early on in the pandemic, was more for the rest of us. There wasn't actually copious guidance, now that I read this properly, for care homes. The first guidance that came out from the Scottish Government that I recall for care homes was guidelines on outdoor visits, and that was in the June. But it always took a while, and that was maybe one of the easier bits of guidance for care homes to implement. It was outdoors. But other guidance that came later in the year for care homes, which was -we'll maybe come on to this -- about trying to ease some of the restrictions on visiting, there was very often quite a big delay in that guidance coming out and something happening on the ground that was meaningful.
Q. You talk about Care Home Relatives Scotland. How did you make contact with them?
A. I came across them through another relative in the care home who I'd made friends with. We liaised a lot by phone, social media, and they suggested, "Get on to Facebook. There's a really good group there that's talking about all the issues we've been talking about and you'll find that what we're experiencing is across the country". So that's how it started for me. I just got involved through Facebook and reading everything that was being said and finding there was this huge
common cause and empathy.
Q. Did their website or Facebook page contain lots of information?
A. Yes, always. It would be great. I mean, whether it was giving you a link to a newscast that had been and you could watch it again and see what was being said or whether it was a steer or a heads-up on when there was going to be some -- perhaps some discussion, maybe even at Cabinet Secretary level, what to do, who to lobby, the fact that the group was going to divide into regions to become more focused -- so every step of the way, every day just about, there was something from Care Home Relatives Scotland to help guide me to focus my energy into trying to help -- to help us but also to help the wider cause as well.
Q. So you were being provided with information by Care Home Relatives Scotland. Did you use that information? How did you use that information once you'd got it?
A. So particularly where we were asked to put pressure, political pressure, to try and effect change -- if there was anything, whether it was to sign a petition or to write to an MSP, I would do that, whatever was asked, and I would use the framework of the focus they were trying to make, which was to get meaningful access and eventually to essential care-giver status. But these

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were objectives that were very clearly coming out of the campaign group and, if they asked for help to lobby, then I would be doing that.
Q. Were you in communication at this stage with the care home?
A. I was always in contact with the care home on a sort of -- this business about phoning and asking how she was, so I was - that was at that level, but I wasn't particularly flagging up things that were going on in a national context. When I was dealing with the care home, I was speaking about Mum.
Q. Right. Did you draw to the attention of the care home any of the new guidance as it came out?
A. I did, latterly. It was in October, by which stage it was a frustration, and from October onwards I didn't ever hold back on drawing attention to any guidance that had come out.
Q. You didn't hold back?
A. No.
Q. Really?
A. Politely, persuasively, hopefully.
Q. So you're talking, at paragraph 46, about new guidance coming out on 12 October, but earlier in the same paragraph you had referred to writing in the strongest possible terms. Tell me about that.

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A. So this is -- my paragraph dates are slightly out of
        sync here. On 30 October, that one, I wrote to the
    manager of the care home and I copied in senior
    management, asking when they would implement recently
    updated Scottish Government guidance. Now, that
    guidance came out on 12 October and it was to give care
    home residents more meaningful visits and the guidance
    said it would be four hours of visiting contact in the
    resident's room and with touch. So this was
    a significant change because prior to that there had
    been these outdoor visits which had been dreadful I'm
    sure for many. They were dreadful for us. So to get
    this guidance and to see this guidance was a huge
    relief. But I went in in October -- my sister went in
    very early October and had the most distressful indoor
    visit with our mother in an interview-style room and she
    wrote it all down in an email to me to offload. It was
    appalling. So it was on the back of that experience
    that I wrote in very plain and strong terms on
    30 October to ask when they were going to implement this
    guidance that was allowing for four hours in the
    resident's room with touch.
Q. And that was }18\mathrm{ days after the guidance had been issued?
A. Oh, yes, yes.
Q. Okay. Did you write to anyone else around the same
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time?
A. I wrote to my MSP on 18 October. There was going to be
a discussion, as I understand, at Cabinet Secretary
level, which was an important discussion because it was
to raise the awareness of the importance of the
care-giver in the resident's emotional health and
well - being and also it was to push for a Scottish pilot
scheme about testing a dedicated care home visitor. So
I wrote a letter to Mr Ewing to add his voice in support
of getting these changes made and that recognition --
the important bit was the recognition of the role that
the essential care-giver had in the life of the
resident.
Q. Now, at this point you've fired off two letters, the first one to the care home. What response did you get from them?
A. I didn't get a response at all. I very often would write and fire off my letters, my strong letters -polite, strong letters -- but sometimes a letter like that would eventually result in a change for us, and the change for us, on the back of that letter of 30 October -- so that was at the very end of October -I was given -- and this term started to appear in October -- "essential visits". So I was told I could get an essential visit to our mother's room, which is
what this 12 October guidance was all about anyway, but I was granted. So perhaps my letter of 30 October resulted -- it was a result.
Q. What about from Mr Ewing?
A. Yes --
Q. What response?
A. -- I got a reply, a polite reply, and very concerned for his constituent and very supportive and to say that he would be supporting and would raise it. But it was short and he just referred to another letter -- and he let me have a copy of it -- that had been issued by Cabinet Secretary, but that letter had been written in July, so I just -- you get that feeling of, "Yes, you know. I've asked you a question now. We're in October. This is urgent and you're supportive, you've given me a response". The Government had been discussing this, had recognised the impact that separation was having on people in care, so this conversation had been going since July and we were still trying to get traction and get meaningful visits inside care homes in October.
Q. I' II come back to that phrase, "meaningful visits" -A. Yes.
Q. - - towards the end of asking you questions. But at the moment there was formal guidance on garden and window visits before indoor visits started. Did you have

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## garden or window visits?

A. Window wasn't going to be remotely possible. Our mother could hardly stand. I mean, she did walk into care when we took her in in that June, the nine months previously, but by this stage, with the progression of her conditions and aging and all the rest of it -- and she was walking towards end of life -- she couldn't have got near a window. She was impaired hearing. Visually, if you have somebody outside a window, they're silhouetted --
Q. That wasn't going to work?
A. That wasn't going to work. And the other one was garden. We had a few in the summer. I think all my siblings got a chance to have a garden visit and all found it very distressful because she was on the other side of a large table, under a gazebo, eventually, and we were masked. She couldn't recognise us. She used lip - reading to enhance her hearing. So really these were very distressing for us and her types of visiting.
Q. Okay. In the autumn of 2020 indoor visits start. Now, at the care home your mum was in, did you have indoor visits in the way that they initially operated?
A. We did. In the initial operation --
Q. Tell me about the room in which they took place.
A. Yes, yes. For everyone, visiting this care home, a room
was set aside very near the front of -- the entrance to the building, and it was like a management room, small, office -style. It had one seat like the one I'm sitting on for me and Mother was brought along in a wheelchair at the appointed time, and there was yellow tape across the floor at 2 metres' distance, "You must not cross", and you were given instructions. Masks, gloves, pinny, and half an hour.

This was beyond distressful because our poor mum had had to be hoisted out of wherever she was comfortably, bed maybe, possibly at her recliner chair -- that's all she was doing at that point -- put in a wheelchair, wheeled along to a room and then wouldn't even recognise the person sitting there because we were masked up and she was seeing people in masks all day long.

So we -- part of the reason I wrote on 30 October, without threatening it -- but the point I was making was we ourselves were becoming the agents of distress for our mother and we were not prepared to do that any longer. So without saying it, it was almost like, "Unless we can get to see her in the comfort of her own room, we will not be putting her through this again".
Q. Could you read the final two sentences of paragraph 53?
A. From "They sat ..."?
Q. No, "It was a room ...", the final two sentences.

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A. "It was a room with a glass door and watched over from the outside by a staff member to ensure there was no effort to touch or move closer. Could a room or situation be any more inhospitable and hostile?"
Q. When was this?
A. That was 4 October.
Q. So that was at the time of Eat Out to Help Out?
A. Yes, I think so. Yes.
Q. Paragraph 55, what you say there is that any guidance given out by Public Health Scotland was immediately trumped by a single positive COVID result, often from a worker who might have picked it up outside. Did that have -- how did that interfere with building a relationship -- rebuilding the relationship between yourselves and your mum?
A. Well, we didn't see her very often because of the cycle of long gaps between it being possible to go in and see her and you were thinking you might get to see, but once a week for an hour, it wasn't giving her the amount of contact she would have needed anyway to retain that -meaningful contact was all about nurturing her. The whole of her was dependent on a reason to live. And so, when we were allowed in and then this would be consistently taken away from us, all that was doing was disrupting an already crumbling relationship we had with
her. We were losing it. We were actually losing the relationship with her. But despite all this, she would still recognise us whenever we did manage to get close enough.
Q. You talk about progressing into winter and your mum's health deteriorating and that being difficult. Can you tell us about that?
A. So we hadn't seen her properly really ever since the start of the pandemic apart from a few garden outdoor visits and a couple of disastrous indoor visits in that interview room. We were given an in-house report on her welfare. The care home did it. They had to do these, and this was one they had to do in our absence and also in the absence of her social worker. In that we noted things like her BMI, her weight loss, and they'd talk about her cognitive decline as well, the fact that two carers are needed at all times to move her. So this was October that was issued -- it was 12 October, the same day as the first meaningful Government guidance on getting people into care homes --
Q. I' II just help Lord Brailsford. This is spoken about at paragraph 63.
A. Yes. So -- sorry.
Q. So you're saying --
A. Yes. The in-house report. It was very clear to us. We

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didn't need that report to tell us that she was on a decline. We could see from the little visits we had that she was on a spiral down. But this welfare report certainly consolidated it and summarised it for us and we took huge, huge exception to this report.
Q. Why?
A. Because it started with saying she enjoys her garden visits and I think, in October, we'd managed to have -her piper grandson came and played outside the grounds, outside the communal window for her, and "enjoyed seeing her family and enjoyed hearing the pipes", and yet my sister 's account of it and my own account and my brother's account were that these meetings in the garden, outdoors, and the ones in her room in October were so distressful for her that we were not going to do them any longer. We couldn't put her through it. So you had this huge "Yes, it's all alright" in the report and the reality.
Q. In days of yore, when we were younger, we used to read or see reports about Pravda, the Russian newspaper, and the front page was always, "Tractor production is up". It would always put a positive spin on everything.
A. Yes.
Q. Was that what this was?
A. An unbelievably positive spin. There was a small bit
that said "Relatives' comments", and of course that was blank because we hadn't been consulted on this. So I wrote what I thought was the situation about her decline in health, visible decline in health, and all the evidence is there in terms of the technical, the factual stuff --
Q. Was that written in terms which were polite but forceful?
A. That was another polite but forceful but getting more forceful than polite. No, I was never rude.
Q. No. I'm going to ask you to read a couple of
paragraphs -- I hope this isn't difficult for you -- but it 's paragraphs 58 and 59.
A. Okay.
"For Christmas Day 2020, we asked if she could be brought in a wheelchair specifically to the large window of the communal area of her wing. The carers obliged and did their best to help her understand who was outside. This was a repeat of an approach that we had trialled in October with each family member holding a placard with large letters forming their name and held it to our chests. The carers read the names out to her as we approached the glass.
"We all took turns of coming up to the window, in our family groups, and a grandson played some stirring

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reels on his bagpipes which she surely heard. The great-grandchildren danced. I pressed my face close to the window, I saw her tear-stained face and I could see her raise a finger and point, weeping and saying to the carer inside, 'that's my family'."
Q. It continues:
"Our time was up, we had to leave the grounds after a bit, but we didn't get inside on this her last ever Christmas, we had opted to all be there and to try to be 'with her' from the outside. It was harrowing to see her like this, separated by glass from her loved ones, no possibility to speak or hug. Was this a better option for her than seeing just one of us indoors? We will never know. It was probably marginally better for us all to be together, seeing her from the outside, but it was nonetheless a very emotional ... day."
A. Yes.
Q. I think we can understand that.

At paragraph 62 you talk about the downwards spiral of your mother's health and you say at the end of paragraph 62:
"We firmly believe lack of meaningful contact with her loved ones was having a profound effect on Mum's mental and physical health \& well-being."

Tell us about that.
A. I think from what we could see of her through the window at Christmas, from what we read about -- from the internal report and from the three visits that -I managed to get three visits indoors from November up until 15 December, and this was on the back of my strong letter about "We need to get in", and I was granted essential visitor status, so --
Q. How did you find out about essential visitor status?
A. I only heard of it through Care Home Relatives Scotland. This again was another -- the fountain of all good information for us to actually use as a tool to try and make the case we needed to make with our care home.
Q. And you didn't get it from the care home?
A. No, no.
Q. And you didn't get it from the Scottish Government?
A. I think eventually you're hearing words through Scottish Government guidelines but maybe not until about the February of 2021. But back in October, after I'd written, after the guidance came out saying we should get more meaningful visits of four hours with touch -and I pressed to get in and I used the words because I'd picked it up from somebody else, and somebody else had heard it from Care Home Relatives Scotland. I heard it too --I used that term almost as if it was a right for us or for her actually to have a relative going in with

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an essential visit. So I had three essential visits, if you like, some time in November to about 15 December, but only three, and it was just once a week, which really was not ever going to be sufficient for our mother.
Q. And were they timed? I mean, were you told start at 2.00 - -
A. Yes. Yes. It was an appointed time and for an hour, which again wasn't helpful, and I wrote at another point to say, "Great. Thank you so much for allowing this", always very positive, "Thank your staff for how much they've helped to facilitate this, but ...", and the "but" was, you know, the time of day that I was appointed to go might not have been a good time for her.
Q. Tell me about that.
A. So she might have been asleep and very asleep by this stage, semiconscious, that kind of deep sleep, so if I was to go in and see her, she might not have been all that alert or awake, but other times of the day she might have been having a better time and the carers did say, "Yes, your mum is sometimes more alert and in better form at other times of the day". It's almost like, "It's a pity you couldn't have been here then". It's kind of like, "Well, that's the problem. You've given me an appointed time on an appointed day. It may ideal world it would have been great if a carer could have seen that she was in good form that morning and got on the phone to me and say, "Can you get here in half an hour?". That would have been ideal, but it didn't work like that.
Q. You then move on, at paragraph 69, to talk about your mum becoming very weak. Then at paragraph $71--$ sorry, give me just a second. Yes, 70, you talk about essential visits for care homes frequently mentioned in the Government agenda. Although it was mentioned in the Government agenda, could you see real progress in that regard?
A. No, I didn't, and this is actually a very important point. I had managed to get granted essential visitor status in November but only got three visits, but that was better than had been. On 15 December the care home asked us what arrangements we were wanting to make for Christmas Day because there was going to be a huge demand and there was an awful lot of things to factor into that and they couldn't have everybody in the care

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home, so I - - we, as a family, opted to say, "Look, we've actually at least had essential visits for a bit of November and a bit of December. Let's get Christmas and New Year over with and we'll pick it up again in January. We will do instead this outdoor at the window thing", which we did.
Q. With the piper?
A. With the piper, we did.

But on 23 December we got an email from the managing director of the care home to say that as of the 26 th, so from Boxing Day, the whole country was moving into Tier 4 and that meant visiting with loved ones in care homes should be limited to essential indoor visits only. And that was further qualified by saying essential indoor visits only would only relate to people at end of -- who are needing end-of-life care or experiencing extreme stress.
Q. Right. Let's unpack that a little bit. You had been granted the status of essential visitor back in November?
A. Yes.
Q. Were you ever told what the definition of "essential visitor" was?
A. I wasn't ever told it. I think my own understanding was that everybody was getting or trying to get an indoor
visit and that was generally in this interview room, so to get an essential visit was what we should have been getting anyway because that's what the 12 October guidance was suggesting, but an essential visit was into the resident's own room and it was time-limited and it was all caveated with all the usual stuff around PPE and not hugging. But essential visitor status, in my interpretation of what I was allowed, although I was never told this -- I never saw a policy, but I was -- my understanding was that I would get into our mother's room for that visit to prevent her having to be hoisted and wheeled to somewhere else. Does that make sense?
Q. Hmm-hmm. Were you ever given a definition of what was meant by "end of life"?
A. No, never, and I only thought to ask that when it was clear that that's where we were at. And it became clear to me - - even in October I was thinking - - looking at her frailty, her weight loss, her general cognitive skills, the fact that she had to be fed in tiny little pieces, I knew that we were somewhere on the journey to end of life, but it's defining the indefinable. It's very hard to know when somebody is going to be at -- and I have no experience of that from previous. Clinicians probably do, but, for me, we didn't know. We hoped we would be told. We were all aware it was nigh.

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THE CHAIR: Actually the words "end of life" and indeed, for that matter, the word "essential" in the context of visitor status are both -- would you agree? -- entirely subjective?
A. Yes.

THE CHAIR: And you're very correct that you have no experience of medical matters and therefore your view of end of life is probably even more subjective than a clinician's view.
A. Yes.

THE CHAIR: And "essential" applies to everyone. You would have thought that it would have been helpful if an attempt had been made to define these terms. Now, I fully accept that an attempt to define those terms is itself wrought with difficulties, one that I might unfortunately have to wrestle with in a period of time after I've heard all this evidence. But do you agree that the subjective nature of the terms makes it probably desirable that some effort should have been made to define them in a way that at least would have been consistent?
A. I would have so welcomed that and I did ask actually several times once we got into January because I saw that that -- "end of life" was almost a qualifier for getting essential visitor status.

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MR CASKIE: The other qualifier for getting that status was
    extreme distress?
A. Yes.
Q. Was that defined, as far as you were aware?
A. Not to me, and -- I was aware of that with another
    resident actually and I was aware that they got
    essential visits on the basis of extreme distress, and
    our mother was not in that category of extreme distress.
    She would have been in extreme distress when hoisted and
    that then put her into a frame of mind and disposition
    that we could not have a meaningful engagement with her,
    no matter, but she wasn't frequently experiencing, as
    far as we know, the sort of distress one might imagine
    that refers to.
Q. So that -- arising from what Lord Brailsford has asked
    you, that's another subjective term?
A. Yes.
Q. Except it wouldn't be a subjective term if someone had
    defined it?
THE CHAIR: Well, it might have been subjective but at least
        there would be a basis specified by someone who had
        authority to make such a definition, and that would have
        been of some - I I think to be fair, you're saying that
        that would have been of some assistance.
A. It would have helped.
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THE CHAIR: You might have cavilled against the definition, but theoretically you would have had a chance to challenge that, perhaps, but nevertheless it would have been something.
A. Yes, and there was nothing.

MR CASKIE: And the other group that it would have helped was care homes --
A. Yes.
Q. - - because they presumably were struggling with those definitions as well?
A. Yes.

THE CHAIR: Well, actually, in fairness, I now see, since it 's come up on my screen, you wrestle with that problem at paragraph 78 or you attempt to, so you're well aware of it.
A. Yes.

THE CHAIR: I see also you draw my attention to a number of other highly subjective statements, "extenuating circumstances" and so forth. It's all entirely subjective.
A. Well, that "extenuating circumstances" came up on the letter -- the email that came out on the day of lockdown, very short, to say, "We will be locking down but we will of course [it said, something like that] allow visits under extenuating circumstances". Now at

MR CASKIE: Can I take you to paragraph 77 and ask you if there's anything that you want to say about what's said there?
A. Do you want me just to read it?
Q. No, I want you to tell -- Lord Brailsford is capable of reading it, so do you have anything to add to it usefully or is it simply an effective stand-alone paragraph?
A. I think in some ways we have covered it in that the care home wasn't prepared for defining "end of life" or "essential visits" and it surprises me in a way because that is their job. This is what they do and this was now almost a year into the pandemic. They must have had

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many, many people at the end of their life and passing away and they must have had to grapple with that. So it 's just perplexing that this hadn't been brokered earlier.
Q. Now, the next things that you provide are details of what you head "Closing stages".
A. Yes.
Q. Can you read the second sentence in paragraph 81?
A. "I was granted 'unlimited' access from that point, and I visited every second day for the next 11 days."

## Keep going?

Q. $\mathrm{Hmm}-\mathrm{hmm}$.
A. "Even at that late stage in her life, she recognised me and while she couldn't talk much, what she did say really surprised me. At times she was lucid and appreciative and reflected with great fondness on her family."
Q. Any other relatives visiting at this point?
A. No. It was me. This unlimited access was granted to me on 26 February, although I'd been given -- we'd been given a heads-up two weeks earlier, round about the middle of February, that they were concerned that she really wasn't eating very much or drinking much and that they would monitor it. So we were pretty frantic by that stage, particularly because, although I had been
given essential visitor status back in October/November and had gone for the voluntary break over the festive period, when it came to early January and I tried to re-establish my essential visitor entitlement, if you want to call it that, I got an email saying that, because of the levels of COVID circulating in the community, that was being suspended.

My essential visiting status was suspended and effectively I hadn't been in there since 15 December.
The reason it was was in early January, in our community, there was a sufficient level of COVID circulating that meant Public Health Scotland probably had given a direction about suspending all visits, and that included me, so I didn't really have any special status at all. My visits were just no greater value than suspending people who were visiting in the interview room, so I didn't get in in early January.

Beyond that I was told, "The COVID vaccination is coming soon to our care home. Hopefully, once we get everybody vaccinated, we will be able to relax things again". But that, again, took another few weeks, and so we got to the stage here in February where I'd been alerted to say, "She ain't eating much", and there was a few frantic emails then to say, "When are you going to let us in?". So we were at end of life and I still

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wasn't getting in for all these other drivers out there.
Q. You talk next about effectively a window visit by
a number of family members --
A. Yes.
Q. -- and you bring the piper again.
A. Yes, the piper. Do you want me to read anything there or --
Q. Well, just give us a summary of it. What happened?
A. So it was 19 March and I had managed to get my visiting as an essential visitor reinstated when I think they must have recognised this was end of life, and I was going in, as I said, every second day for about 11 days, and I reached a bit of a burnout with that emotional burden, but also very conscious that my siblings had not seen their mother pretty much for a whole year. I mean, some very fraught and distressing outdoor visits and a couple of room visits that were awful, but they hadn't seen her. And because we knew --I knew, we all knew -this was end game, I pleaded with the care home to let my two siblings, who had seen least of their mother, take my place for a few -- a visit, and they agreed to that. So a brother and my sister took over from me. So it wasn't as well as me, it was instead of me, and they got a visit each --
Q. Not simultaneously?
A. Not simultaneously -- or did they get one where they were allowed in together? I can't remember what the rules were then around families who weren't - - you know there was all bubbles things. That played into this as well. But they each got a visit and we agreed with the care home that, you know, for the time being we would continue with that to give them a bit more time. And my elder brother had stepped back to let them have it because he was the one that lived closer and had seen just a little bit more. So that's where we were at with it.

Then on that day, that 19 March, my sister was - - it was going to be her turn to come up, one of two visits in the week, and she couldn't come because she was unwell from her COVID vaccination so I stepped in to do that visit. And I'd arranged for one of the grandson pipers to come and play again at the window, like we'd done before, which was all organised and agreed and set up, and I was so conscious from my other brother that this was very close to the end that I encouraged him to come along with his wife and his daughter and her partner and the two great-grandkids just so stand beside the piper, to give the piper a little bit of support, to give me a bit of support on the inside and to see his mother for the last time.

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Q. And what time of day did that happen?
A. 2 o'clock in the afternoon.
Q. When did your mum die?
A. Well, she died on the Monday. But when we were outside
her room, the piper started, and somebody from the care
home came in and told me that these people hadn't been
authorised to be at the window, so I was --
Q. Was the window open?
A. No, no. She was lying in her bed well away from the
window and she was dying at this stage. But I was told
to tell them to leave, so that's what I had to do.
I had to say to them, "You've been asked to leave", and
they had to leave the premises. They were ordered off
the premises because that had not been part of the plan
or hadn't been approved by somebody senior enough.
Q. And was there no one there who could just take
a decision there and then?
A. No, no. The manager of the care home didn't work on
a Friday, so the person who was left to assess what was
going on here, all they saw was, "This hasn't been
approved. These people have to go. They're not
authorised to be here".
Q. You say at paragraph 92 that you weren't happy about the
end-of-life plan. Can you say a bit more about that?
A. Sorry, I m not quite sure what you're asking me to ...
Q
Q. And what time of day did that happen?
A. 2 o'clock in the afternoon.
Q. When did your mum die?
A. Well, she died on the Monday. But when we were outside her room, the piper started, and somebody from the care home came in and told me that these people hadn't been authorised to be at the window, so I was --
Q. Was the window open?
A. No, no. She was lying in her bed well away from the window and she was dying at this stage. But I was told to tell them to leave, so that's what I had to do. I had to say to them, "You've been asked to leave", and had to leave the premises. They were ordered off the premises because that had not been part of the plan or hadn't been approved by somebody senior enough.
Q. And was there no one there who could just take a decision there and then?
A. No, no. The manager of the care home didn't work on a Friday, so the person who was left to assess what was going on here, all they saw was, "This hasn't been approved. These people have to go. They're not authorised to be here".
Q. You say at paragraph 92 that you weren't happy about the
A. Sorry, I'm not quite sure what you're asking me to ...

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Q. Sorry.
THE CHAIR: It's possibly background for me to use
        (inaudible).
MR CASKIE: Okay, that's fine.
THE CHAIR: It seems clear.
MR CASKIE: Let's move on to paragraph -- well, at the end
        of paragraph 98, there's a heading, "Outcomes Sought
        from this Public Inquiry", and you list those. As you
        can see, Lord Brailsford can read that. There are
        a couple of questions that I want to ask you about
        paragraphs 102 and 103 - - could you read paragraph 102?
A. "This fact is so often overlooked in the media ..." --
Q. Sorry, I need to go back --
A. Go back a wee to the fact.
Q. - - to which fact.
A. Yes, what fact.
Q. Aha, about the importance of family.
A. Which paragraph?
Q. Sorry, if you look at 101 --
A. Yes.
Q. - - there's a request to the Scottish Government to
        publish figures, and they're about the number of people
        who died not of COVID --
A. Yes.
Q. -- in care homes.
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A. Yes.
Q. And then you say, "This fact ..."
A. The fact that so many people died in care homes
    throughout the first }12\mathrm{ months of the pandemic --
    I believe the figure is something around 16,000 died.
    300 were dying per week in care homes in Scotland. Not
    all of these people but the vast majority will have died
    from reasons other than COVID and, I think that figure
    should be articulated at some point as a result of this
    Inquiry or to help inform this Inquiry because it's an
    awful lot of people -- right? -- that they were in care
    homes, they had no meaningful contact, and for us it was
    a whole year of no meaningful contact, it was
    meaningless contact, with their loved one, and these
    people died alone and lonely, effectively imprisoned in
    their care home, and that was for the remainder of their
    lives.
Q. And their death was not related to COVID?
A. Yes, and I'd like to know, of the 16,000 people who did
    die in that first year, you know, how many suffered and
    had to endure this, no meaningful contact, and then they
    just passed away.
THE CHAIR: Well, actually all of them because everyone in
        a care home, to a greater or lesser extent, depending on
        the care home - - but they all experienced -- I'll use
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    a non-emotive word for obvious reasons - -
A. Yes.

THE CHAIR: -- they experienced the restrictions that were imposed on them as a result of COVID --
A. Yes.

THE CHAIR: -- and therefore all of them went through that experience.
A. They all went through that experience, yes.

MR CASKIE: At paragraph 106 you make positive comments about Care Home Relatives Scotland. Why was an organisation like Care Home Relatives Scotland needed in your view?
A. It was a voice. There was no voice at all to champion the cruelty around withdrawing meaningful contact from people in residential care, be they young or old. The Care Inspectorate even talk about the fundamental importance of meaningful contact in your life and the withdrawal or removal of that has a significant detrimental effect on your mental health and well-being, and indeed, in many elderly people, the will to leave.
Q. Tell me about Anne's Law.
A. So Anne's Law --
Q. I know what it is.
A. Yes, Anne's Law is something that we must have. We must avoid this ever happening again. And Anne's Law would

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give the resident the right to have one designated care-giver, an essential care-giver -- and that's the important word -- the right to enter the care home and give that essential care in the same way that a staff member would.
Q. Could you read paragraph 112?
A. Paragraph ...?
Q. 112.
A. 112:
"The Scottish Government should provide a definition or articulation of 'meaningful contact' for anyone in care, be they old or young, and should enshrine this in Scot's law alongside Anne's law, or as part of Anne's Law."
Q. And then continue on to 113 .
A. "As a family we have struggled to come to the terms with the loss of our mother in this way, knowing that until the very end she was still with us, thinking about the family and never knowing or understanding why she had lost us. Images that we have re-told for this Public Inquiry still haunt us. Our mother lived a long, happy life, until that very last year when everything changed and we were unable to reach out to her and support her when she needed us most. It is painful and difficult to celebrate an extraordinary life when the emotions around

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    the dire circumstances of her final months, and how she
    died, are still so raw."
Q. You then summarise or state what you consider are the
        key parts of your evidence. I think I've been through
        all of your evidence. Can I ask you to look at certain
        of those paragraphs in particular? 118, 119, 123, 124
        and }125\mathrm{ together, and then 126.
A. Can we scroll back down? Can I read because that's --
    I can read and discuss. }118\mathrm{ first?
Q. Yes. We can go back through that. 118.
A. Right. Yes, this is a point that I really want to come
    across loud and clear. It's about meaningless contact.
    So:
            "[The] combined impact of isolation from family and
        meaningless contact sped up cognitive decline which
        contributed significantly to deterioration in both
        physical and emotional health & well-being."
Q. And then 119 is about Anne's Law, and I don't think that
        adds to the evidence that you've already provided,
        although if I'm wrong, please correct me
A. No, that's it. Anne's Law is essential. The right to
        have a family member enter the care home and provide
        that essential care, whatever form that takes during
        a lockdown, the same way that staff do.
Q. 123 --
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THE CHAIR: Before you get there, stop at 121. You ask --
your view is that meaningful contact should be enshrined
in law. As a result of the discussions -- you giving your evidence and the discussion that has ensued during it, would you like to add other terms to "meaningful contact"? Would you like to revise your statement? You don't need to do it. I'm to some extent teasing you, but it flows out of the discussion we had only a few moments ago about the subjective nature of many terms.
A. Well, yes. Yes, I mean I could sit here --
THE CHAIR: Of course you could.
A. -- and paint a picture of what meaningless contact looked like and I hope I've described meaningless contact.
THE CHAIR: Yes, you have.
A. So we need the corollary of that. We need to have -- we must document what meaningless contact -- never forget what that looked like, but we also must write down what meaningful contact should be, and that would be for perhaps lawmakers too, and to get the right words for it, and it's subjective, as is "essential visiting ", as is "end of life". But all of these -- we have learned the lessons here, we're learning the lessons, but we have to have this better -- as best can be articulated so that the ambiguity can be removed and the doubt and
THE CHAIR: Before you get there, stop at 121. You ask -your view is that meaningful contact should be enshrined in law. As a result of the discussions -- you giving
that people could pick up on this instantly should we be faced with this again.
THE CHAIR: I can assure you these are extremely difficult questions but you've posed them.
MR CASKIE: 123, you've made that clear.
A. Yes.
Q. 124 and 125 , read together, is really what you've been talking about --
A. Yes.
Q. -- about, the need for definitions. And then 126, guidance isn't enough, to summarise that.
A. Yes.

THE CHAIR: Another very difficult question but one that you've put in front of us. Not you alone, I should stress, but the other witnesses as well.
A. Yes.

MR CASKIE: I only have one more question for you, and that's this: do you feel as though you've said all you want to say?
A. Yes, I think I do. I think we've covered all the points and they're difficult and very hard to go through, but I'm just very grateful to be able to help in any small way, to give voice to the many thousands of people who haven't had the privilege to have a platform to do this, and I hope that what I'm saying helps in some way to

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articulate what the problems were so that we can review and progress and learn as a society, so that if this ever happens again, we don't make these mistakes.
MR CASKIE: We are the ones who are grateful. Thank you.
A. Thank you, Lord Brailsford, for listening to me.

THE CHAIR: Thank you, Mrs Duncan. The only thing I can say
is that -- you're quite right to pose all these
questions and I'm sure it's meant as help and it is
help, but you make it more obvious how difficult this task is.
A. Thank you.

MR CASKIE: We've got plenty of people to help us.
THE CHAIR: Thank you, all. Tomorrow morning, 10 o'clock.
MR CASKIE: Yes, but not me!
(12.45 pm)
(The hearing adjourned until
Thursday, 16 November 2023 at 10.00 am)

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