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Sir—Richard Horton¹ states that doctors are notoriously protective of their power and prestige. It is possible to think of a nurse who is enabled to prescribe drugs as someone who is nearer a doctor than his or her non-prescribing counterparts, and, thus, as someone who has been promoted. However, a nurse who takes an advanced degree in nursing does not become more like a doctor but becomes instead a better nurse.

The two professions complement each other, but are almost totally different: I would be as uneasy at seeing an experienced and well-trained nurse prescribing drugs for a member of my family as I would at seeing an eminent doctor tending to the same patient's needs of daily living.

Nurses have the complex tasks of observing and assessing patients, identifying problems, and ensuring that individual patients are placed in the best position to do well, irrespective of the specific illnesses with which they are faced. Once patients have been diagnosed and treated, nurses should expedite and coordinate procedures so that the diagnostic and therapeutic activities are completed in the optimum way and the patients' dignity and comfort are maintained.

We nurses think that we do these things better than once was the case, but we have much room for improvement. It is on becoming better at our professional tasks that we should concentrate, rather than attempting to divert our energies into activities such as prescribing drugs.

Instead of comforting sick and troubled patients and ameliorating the environment so that wellbeing and resilience are not compromised, it would be tempting for a nurse with prescribing privileges to offer a drug and depart. In this way the worst among us might truly come to resemble the doctors.

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Aplastic anaemia and telomerase RNA mutations

Sir—Tom Vulliamy and colleagues (June 22, p 2168)¹ report the association between mutations in the telomerase RNA gene (*hTR*) and aplastic anaemia.

In a previous study by these authors,² a rare autosomal dominant type of dyskeratosis congenita (an inherited type of aplastic anaemia) was associated with very short telomeres and mutations in the *hTR* gene. Patients with aplastic anaemia also present shorter telomere length than age-matched controls.³

We have assessed 42 unselected patients with acquired aplastic anaemia (aged 2–57 years, median 26 years) for mutations in the *hTR* gene. Of these patients, ten were classified as having moderate aplastic anaemia, 20 as having severe aplastic anaemia, and 12 with very severe aplastic anaemia. In 17 patients, aplastic anaemia was attributed to drug or chemical exposure, two to post-hepatitis aplastic anaemia, and in 12, a paroxysmal nocturnal haemoglobinuria (PNH) clone was detected by flow cytometry.

None of the three *hTR* mutations described by Vulliamy and colleagues¹ was noted in our series. This discrepancy is intriguing. However, as pointed out by the authors, their study was done in a tertiary centre, and the patients' population could be skewed and not representative of the general population. By contrast, our study was done in patients from one institution, which is the only referral centre for the treatment of aplastic anaemia in an area covering five million of the population of southeast Brazil. Moreover, our patients' population included variable disease presentation, such as drug-induced, hepatitis-associated, and PNH-associated aplastic anaemia.

In congenital aplastic anaemia, especially dyskeratosis congenita, short telomere length has a role in the pathophysiology of the disease, and some cases are linked to *hTR* gene mutations.^{1,2} However, although telomeres are also shorter in acquired aplastic anaemia, our data indicate that mutations in the *hTR* gene are not involved in the pathogenesis of the disease, in which bone-marrow failure is the result of immune-mediated destruction of haematopoiesis.⁴

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Time to do the right things right: Nigerian health care

Sir—The June 8 Reportage from Sally Hargreaves¹ is most welcome. As a Nigerian who trained at the nation's oldest medical school (formerly University College London, Ibadan Campus; now, University of Ibadan), I could not help identifying with most of her observations.

During my own stint at the Lagos University Teaching Hospital, I came face to face with the appalling man-made decay that had befallen the Nigerian health-care system. I was prompted to pursue postgraduate studies in health services and systems research.

As a country that accounts for about 50% of the west African economy,² Nigeria cannot afford to drown the rest of the region.³ Bearing in mind that health and wealth are highly correlated, it becomes urgent that Nigeria addresses its abysmal health and social services to jump start its non-petroleum dependent economy and create a robust society of healthy people who can contribute to nation building.

Like the rest of Africa, Nigeria presents a fertile ground for research⁴ and seemingly miraculous interventions if the right initiatives are taken in a coordinated way. However, currently, there are no up-to-date or accurate data on the state of the health of Nigerians. Research and interventions are, therefore, made difficult. This situation complicates and is further confounded by the near chaotic, inconsistent, foreign aids and programmes aimed at tackling the difficulties.

The Nigerian leadership must rise to the occasion and get its priorities right: restoration of public confidence; increase in investment in health, education, research, and related sectors; creation of sound health and demographic databases; proper needs assessment; the abandoning of inappropriate projects for the identified needs; stimulation of a development of a