

The Health Justice Initiative's 17 Questions on the NHI and Medicine Access

- What specific measures are envisaged to **enable and promote public transparency** related to medicine selection, procurement and contracting processes under the NHI?
- How will the **deliberations of NHI ministerial advisory committees** be made open to the public, and how (and how often) will this information be shared?
- Not everyone living in SA will be regarded as a 'user' of the NHI Fund. In these circumstances, will medicine manufacturers be permitted to sell medicines to health providers other than the State if so, how will this be done, and what will be the maximum price?
- Which medicines will automatically be covered under NHI benefits as part of the NHI Formulary? How will this information be communicated to the public, and how can the inclusion or exclusion of specific medicines be appealed?
- Which medicines will *not* be covered under NHI benefits as part of the NHI Formulary? On what basis will this exclusion be decided, how will poorer patients access life-saving medicines that are not included, and will humanitarian organisations be permitted to access medicines on individuals' behalf?
- Will wealthier people be able to bypass NHI selection and purchase more expensive life-saving and other medicines on their own/with others where the State does not procure these itself?
- Many foreign migrants and South Africans without documentation will not be able to register as NHI Users. How will people who are not registered NHI Users (for any reason) be able to access basic health care services?
- How will the price of medicines not included in or covered by the NHI be regulated? And what role will External Reference Pricing (ERP) methodology play in the NHI and beyond?
- 9 How will the NHI Fund (e.g., the Office of Health Products Procurement, the NHI Board) negotiate with global pharmaceutical manufacturers and suppliers in order to procure for the State, and how will that process be transparent and accountable?
- By when and how will the current Single Exit Price (SEP) system that governs private sector medicine acquisition be amended and/or extended, and in what phases of the implementation process will this occur? How will dispensing fees be charged, and how will Section 21 exemptions be managed?
- Why has the jurisdiction of the Competition Commission been excluded, and which other statutory body will be legally tasked with providing the necessary regulation of price and competition?
- How will the Minister determine that the NHI is 'fully implemented', and what will take place in terms of what medical schemes can and cannot offer members during the transition period, and after the (undefined) date?
- How will the current [medical scheme] provisions related to 'late-joiner penalties', 'waiting periods' and top-up insurance products be managed or transitioned under the NHI without prejudicing existing and also new members?
- Has consideration been given to **designing a competitive and different single medicine pricing system** for SA?
- What specific measures if any will the **NHI Fund** be permitted to take or recommend in respect of **reducing** medicine, to give effect to the intent of 'strategic purchasing' as referenced in the NHI Bill?
- How will **SA's Intellectual Property strategy need to adapt to complement the NHI objectives** of securing the 'best available medicines' in the most affordable way?
- What system will govern how the South African National Defence Force (SANDF) and State Security Agency (SSA) select, procure, and pay for medicines (they are exempt from the NHI), and how will pricing be monitored and regulated under that parallel procurement system?