

Agenda





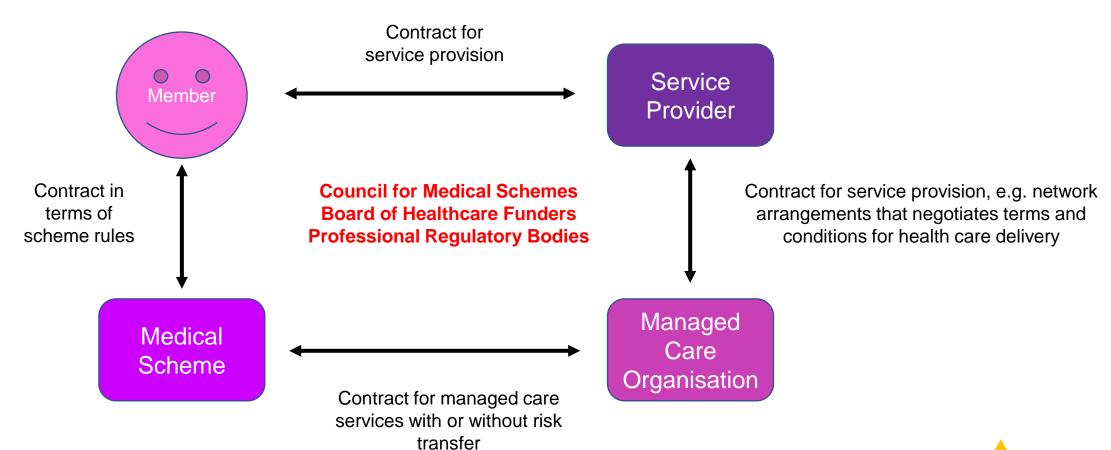
Definition of managed care

Regulation 15 of the Medical Schemes Acts 131 of 1998:

Managed care – is defined as "clinical and financial risk assessment and management of health care, with a view to facilitating appropriateness and cost-effectiveness of relevant health services within the constraints of what is affordable, through the use of rules-based and clinical management-based programmes".

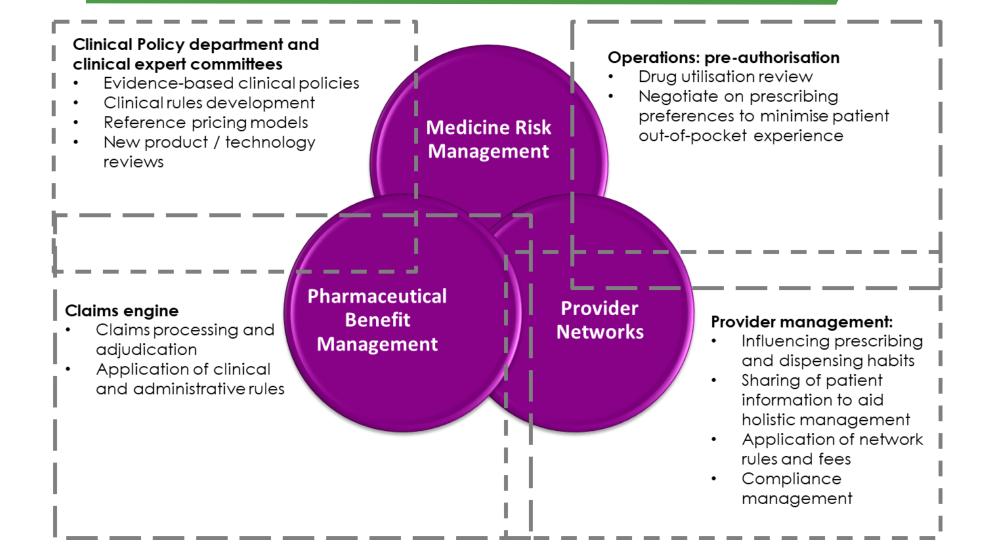


Private healthcare ecosystem





Integrated medicine benefit management





Objectives and operating model

Member and Service Provider Contact Centre Support Digital Health Platforms

IDENTIFICATION

AUTOMATED ALGORITHMS

- Electronic health records
- Acute / Over-thecounter medicine
- Health Risk Assessment
- Disease Risk
 Management
- Hospitalisation

ACCESS

MEDICINE RISK MANAGEMENT

- Pre-authorisation
- Automated disease formularies
- Drug utilisation review
- Benefit design
- Member and provider portals

QUALITY & COST

UTILISATION REVIEW

- Evidence based policies and guidelines
- Formularies/baskets
- Medical Advisory
- Ex-gratia
- Expert panels
- Adverse effect monitoring

UTILISATION

PHARMACEUTICAL BENEFIT MANAGEMENT & PHARMACY NETWORK

- Real -time adjudication
- Clinical rules
- Reference pricing
- Designated service providers

MONITORING & EVALUATION

CENTER OF EXCELLENCE

- Reporting
- Actuarial analysis
- Quality management systems
- Fraud, waste and abuse
- Legal & compliance
- Innovations

Clinical Policy Department
Clinical Expert Committees
Quality Assurance Unit
Risk Management Department



Current gaps in access to medicine

- The existing gap in healthcare access between public and private sector has widened over decades
- Legislative and operational guidelines impacting access:
 - Essential Medicines List (EML) is not inclusive of all the registered products by South African Health Products Regulatory Authority (SAHPRA)
 - Prescribed Minimum Benefit (PMB) algorithms is currently only available for management of Chronic Disease List (CDL) conditions
 - Formularies and guidelines developed by managed care organisations (MCOs) are highly influenced by financial outcomes/affordability
 - Discrepancies in industry in determining reimbursement criteria
 - Single Exit Price (SEP) legislation impacts the availability of products in South Africa and the ability for pricing negotiations to improve access
- Misalignment of incentives between pharmaceutical manufacturers and MCOs



WHO guidelines on pharmaceutical pricing policies

	Pricing Policies	Applied by SA
1	External reference pricing	N
2	Internal reference pricing	Y
3	Value-based pricing	Y
4	Mark-up regulation across the pharmaceutical supply and distribution chain	Y
5	Promoting price transparency	Y
6	Tendering and negotiation	Υ
7	Promoting the use of quality-assured generic and bio-similar medicines	Y
8	Pooled procurement	Υ
9	Cost-plus pricing for setting the price of pharmaceutical products	N
10	Tax exemptions or tax reductions for pharmaceutical products	Y



Way forward

