

ETHICAL ASPECTS ON PRIORITISATION OF NEW CANCER DRUGS

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My background

- Researcher in bio-, medical, health care, public health and health policy ethics
- National council of medical ethics (2000-2006)
- Ethics delegation of the Swedish Society of Medicine
- Member of the pharmacological regional "NICE" of Västra Götaland County Government



Priority-setting and rationing of resources is a core ingredient of medical practice, policy ethics

- A special case of distributive justice: the desirable allocation of available resources in some area. Base question is: according to what principles?
- **Different levels**: society as a whole (between sectors), between areas within a sector, between type-cases within an area, between cases within a type, ...
- Always constrained by (a) higher levels determining the amount of resources, (b) dynamic effects: how a distribution will affect the future demand and amount of resources.
- Basic rule: what you give the one, you deprive another



Basic considerations

- **Benefit to individual patient** (depending on seriousness of disease, effect on disease, side-effects, and evidence-base). Benefits may include QUALY aspects, but also e.g. intergrity and autonomy aspects. **Internal tensions/conflicts may occur**.
- **Benefit to society** (depending on prevalence of disease, level of impairment, likelihood of patients recovering functionality and returning to work, etc).
- **Opportunity costs**: look at all options and patient groups at once and compare effects, not only one option/group at the time this implies a **requirement of cost-efficency**.
- **Normative constraints**, e.g., that patients should never be treated by the harming of other patients or people, that a treatments must respect autonomous decisions of patients, etc.
- **Justice constraints**: that only some of the aspects above should be considered and that they must be considered equally for all (the principle of human worth).
- Swedish law excludes benefits to society from playing any role in priority setting *within* the health care sector, but not *between* health care and other sectors (as in public health management)



New Cancer Drugs: General Considerations

- Very serious diseases, but cancer is as such no more important than other conditions with similar effects (although there is a public culture to that effect: "the Big C").
- Increasingly expensive, either per patient or due to rising number of patients
- Tendency towards more specific strategies ("personalization") → consumer group/ product shrinking → higher prices per unit (pure arithmetic of business)
- **Highly variable and often limited effects**, albeit symptoms may be reduced for a limited time, but usually with quite pronounced side-effects (may still be better than the standard today!)
- **High opportunity cost** as resources might be better used for other serious conditions
- Cost-efficiency goes down → if benefits to society are allowed, this upshot is stronger



New Cancer Drugs: Radical Effect Variation

- In some cases: effect may be relatively good (actaul survival), but also quite bad, this varies in a way impossible to foresee.
- Using such a drug means introducing a lottery through which some patients are probably killed (unintentionally by side-effects), and through this other patients can survive some time more.
- The principle that patients should never be harmed for the sake of someone else seems to apply.
- At the same time, patients may accept these odds, just as they may ask for therapies not considered responsible to use by the profession



The Ethics of Pharmaceutical Pricing

- States/societies must resist the business strategy of steady increased prices of drugs, and this can only happen if they say "no" to some offers, in order to incetivise producer to offer lower prices.
- Otherwise, publicly funded health systems will be systematically milked of resources by commercial companies, and this will hit many more patients much more seriously.
- The price of a drug set by a company is ultimately determined by the financial return expectations of the owners of that company: these are **chosen freely**.
- At the end of the line, these **owners are people who have a moral responsibility** for contributing to a situation where seriously ill patients are denied treatment.



Conclusion

- Given the new strategies of commercial producers of pharmacological products, there are strong arguments for states to resist the offers.
- Also: strong ethical reasons for health systems and health professionals to ration and set limits based on ethically well-founded priority-setting → what this means depends on how much public health perspectives are allowed to play a role.
- Now: nationally centralized price negotiations are coming rapidly, next step is multnational alliances to press prices further, e.g. Nordic Countries, NC + Holland + UK, entire EU (?)
- This is a time of flux, hopefully a new equilibrium will settle itself in the future, where the issue of what drugs are to be publicly funded can be better foreseen.