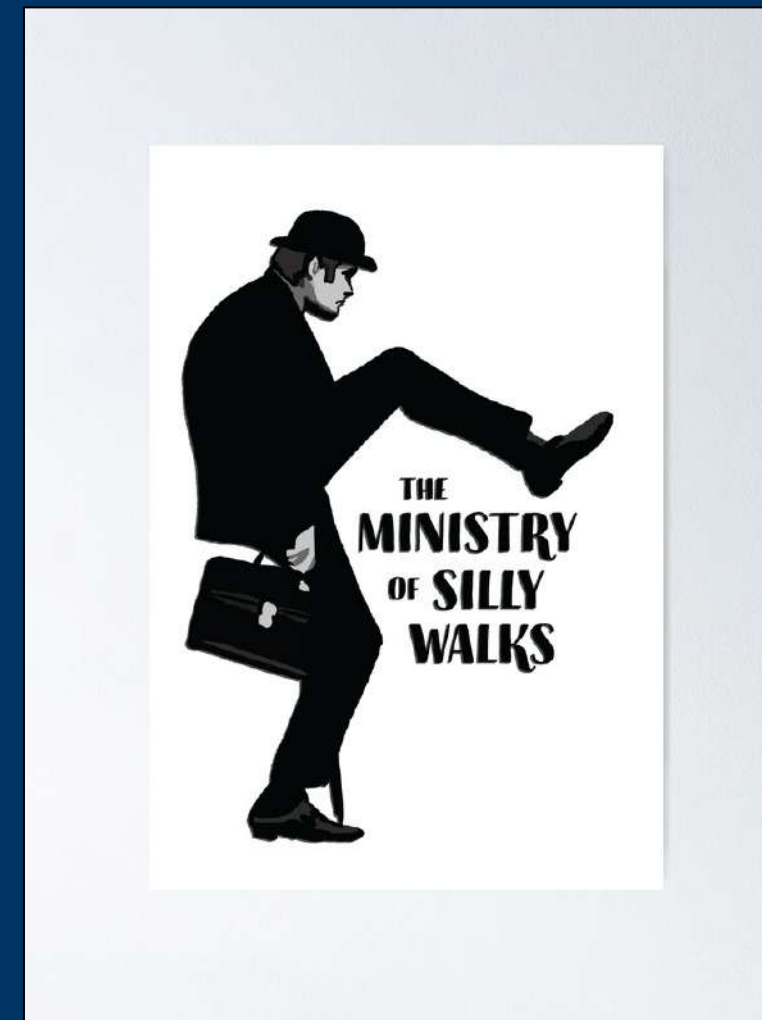




Catalan Clinical Audit
Network for Quality Improvement
in Radiotherapy

What to do when we do not have standards?

Prof. Dr. Dirk Verellen



Co-funded by
the European Union

CAT·ClinART

Objectives of the lecture

- Radiation oncology evolves, standards follow ...
- Some examples
 - MSQA and PSQA
 - BELdART quid?
- Focus on patient safety and quality

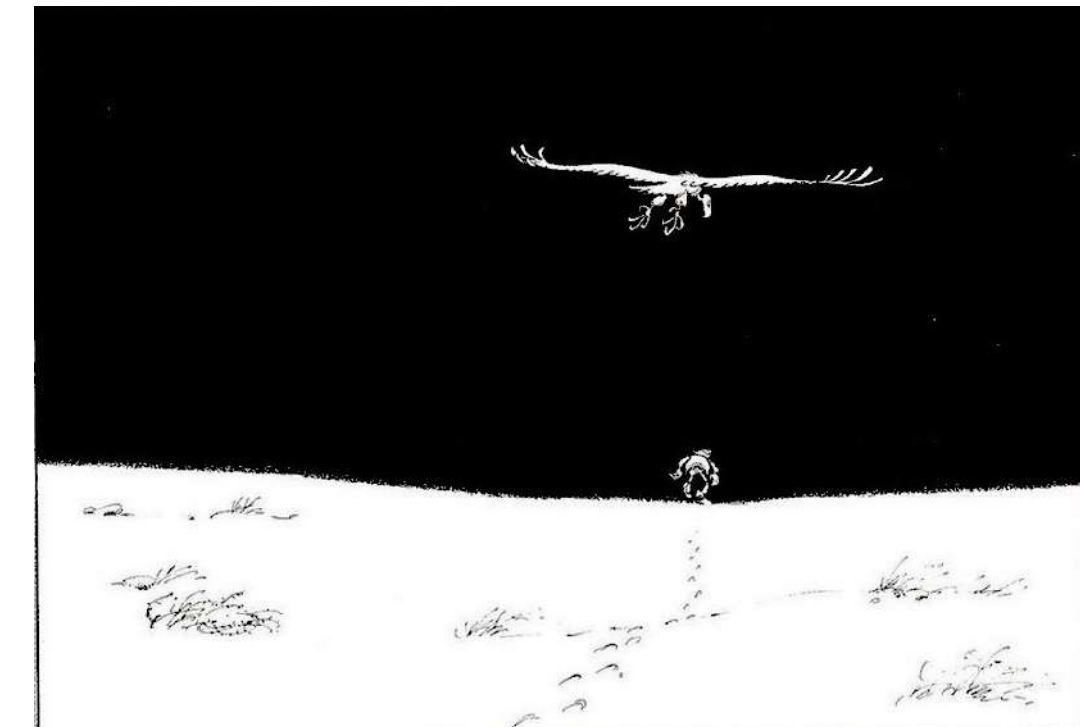
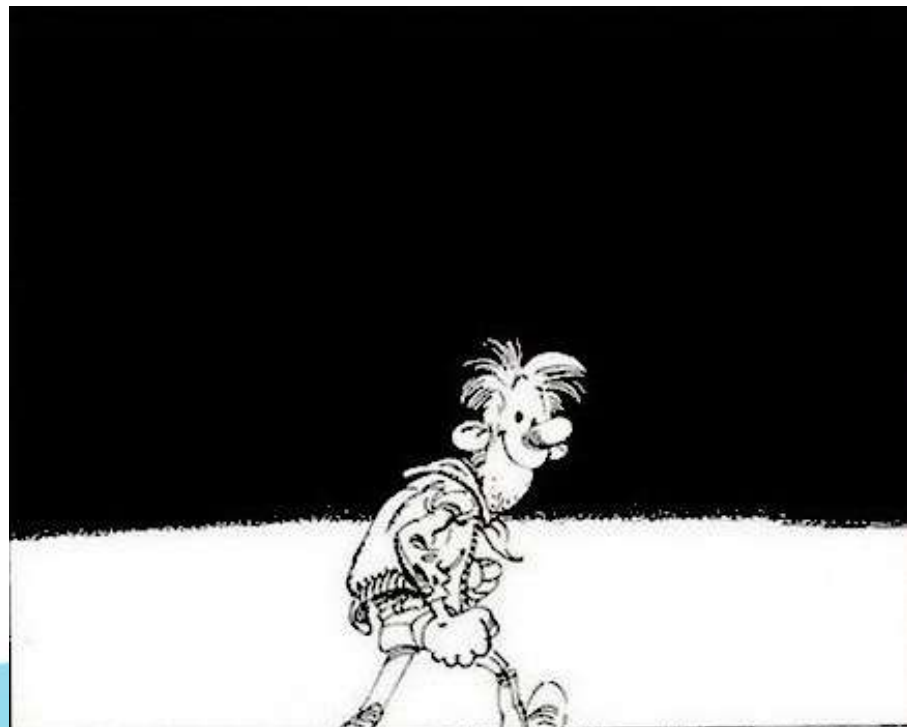


Franquin

IMRT as an example on introducing new technology

Technology implementation races ahead of guidelines

- Small field dosimetry
- In vivo dosimetry
- Independent dose calculation



Franklin

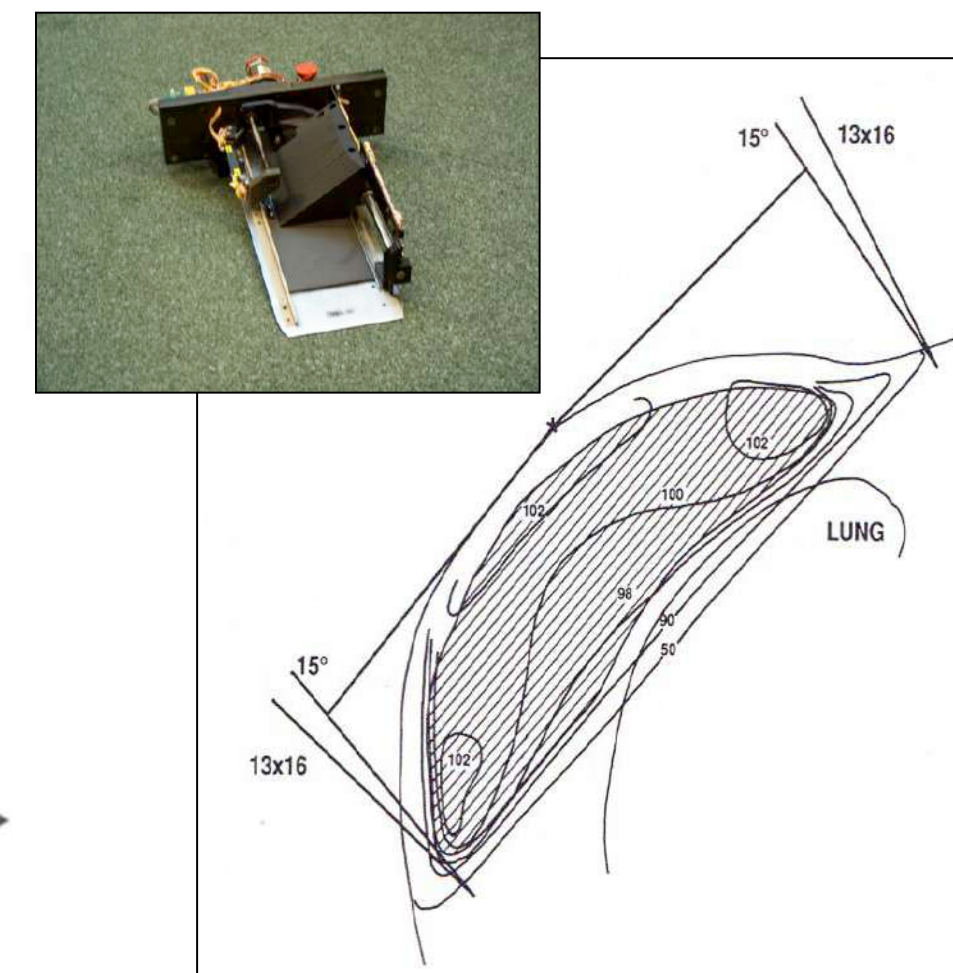
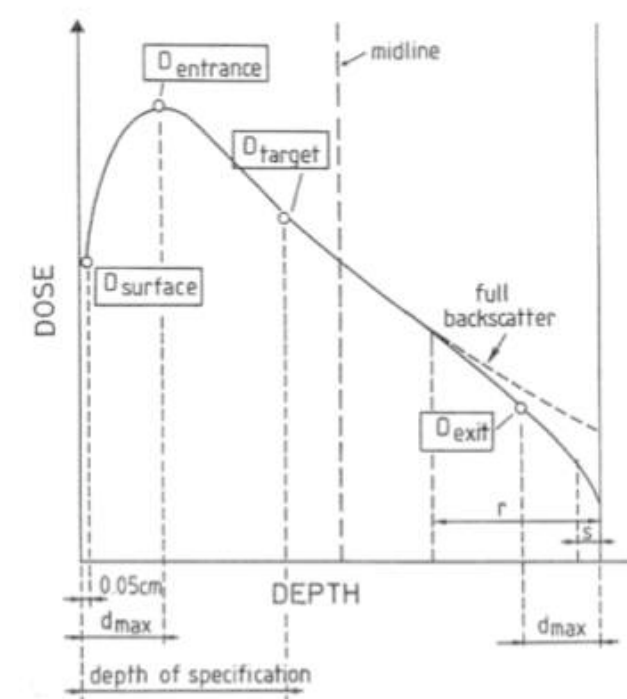
Learning from the past ...

We learn from our mistakes ...

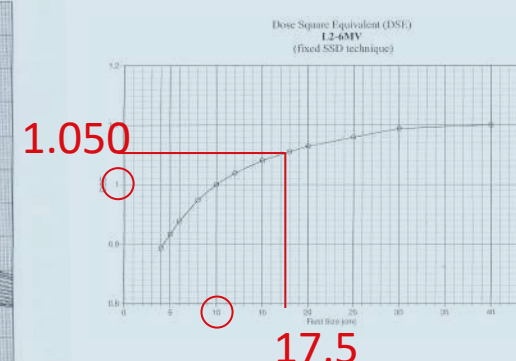
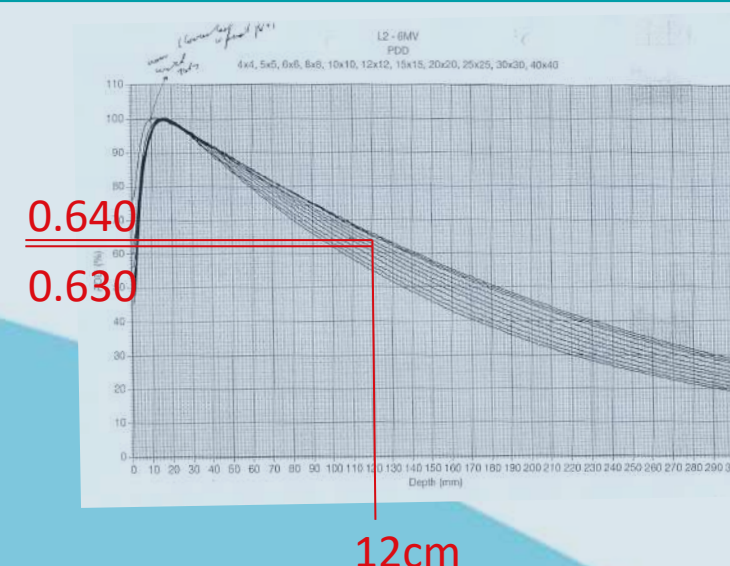
- In vivo dosimetry



- Independent MU calculation



- TD = 200cGy
- d = 12cm
- SSD = 100cm
- FS = 17.5x17.5cm²



$$\text{OF}(17.5 \times 17.5) = 1.050$$

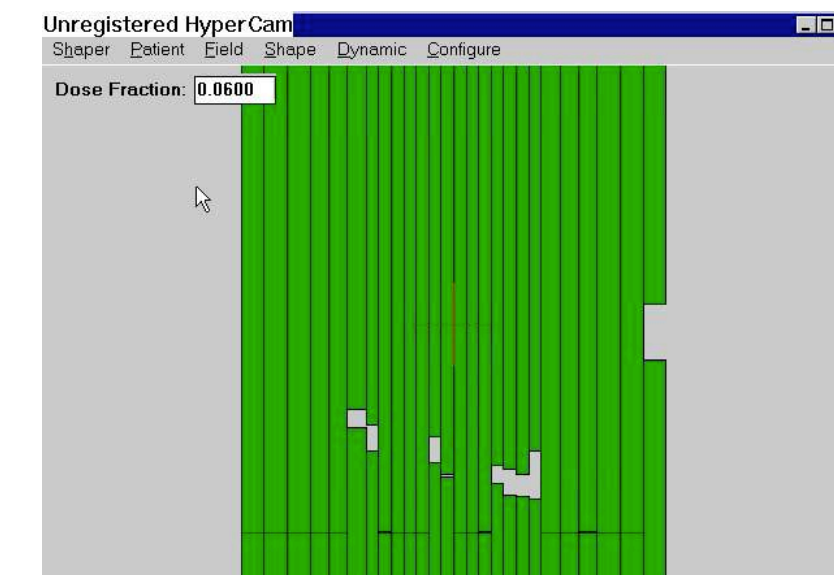
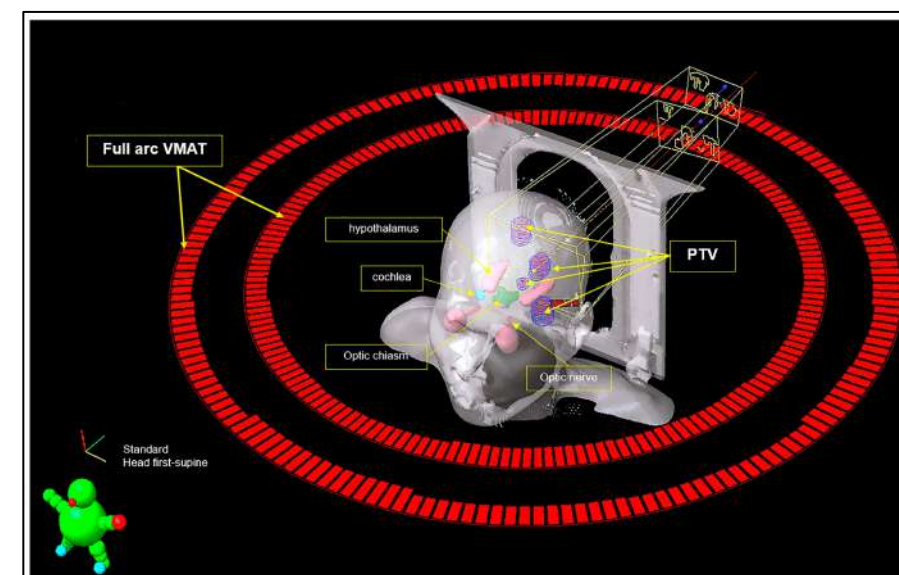
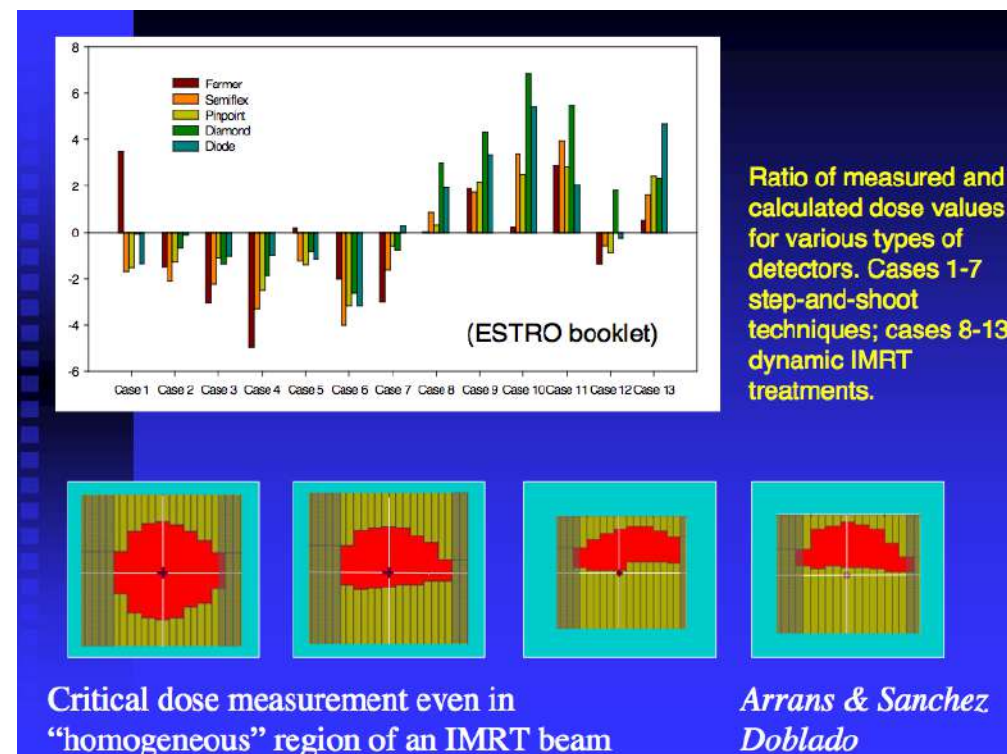
$$\text{PDD}(12, 15 \times 15) = 0.630$$

$$\text{PDD}(12, 20 \times 20) = 0.640$$

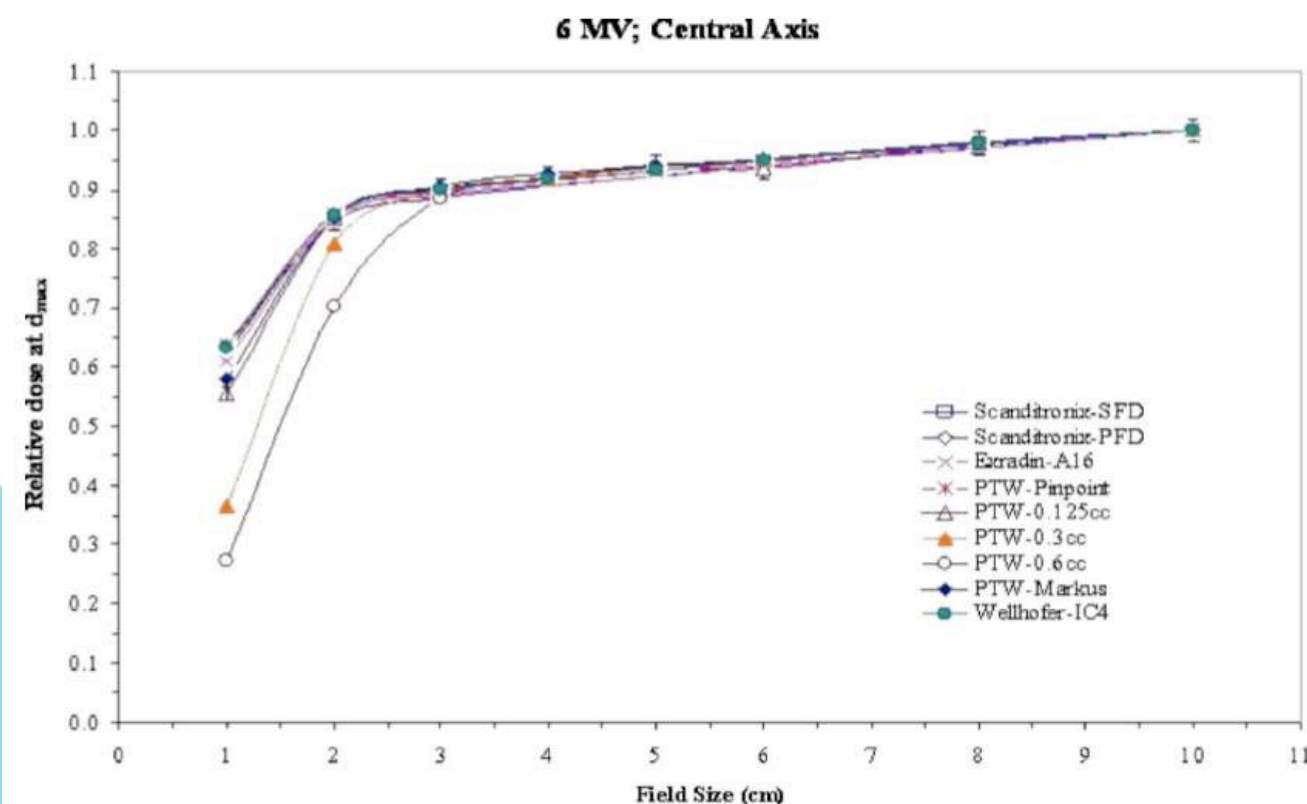
$$\Rightarrow \text{PDD}(12, 17.5 \times 17.5) = 0.635$$

$$\# \text{ MU} = 200 / (0.635 \times 1.050) = 299.96 \sim 300 \text{ MU}$$

Absolute dose calibration



In this case ... size matters



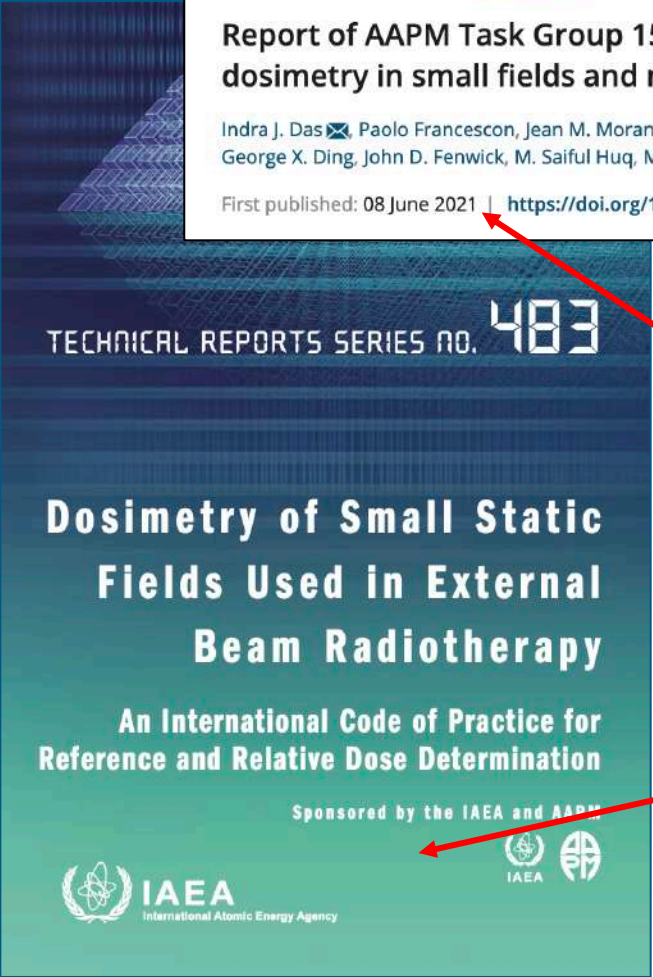
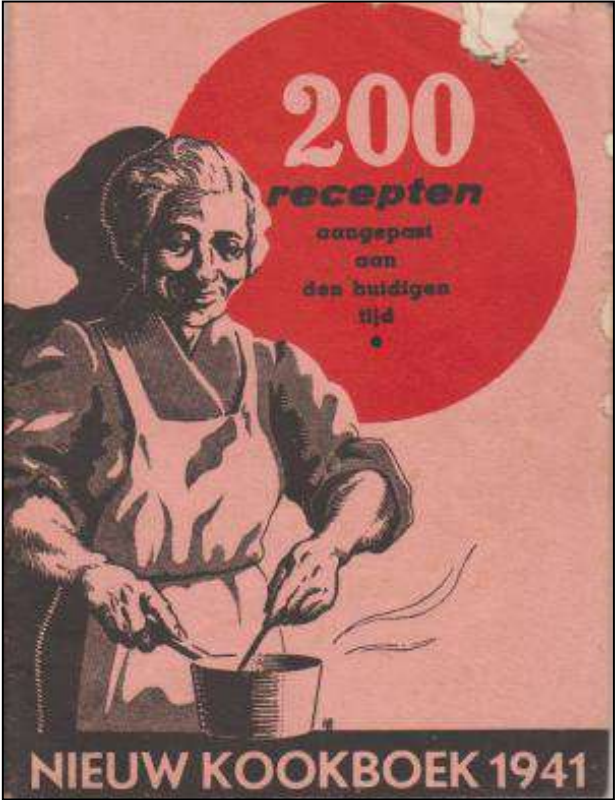
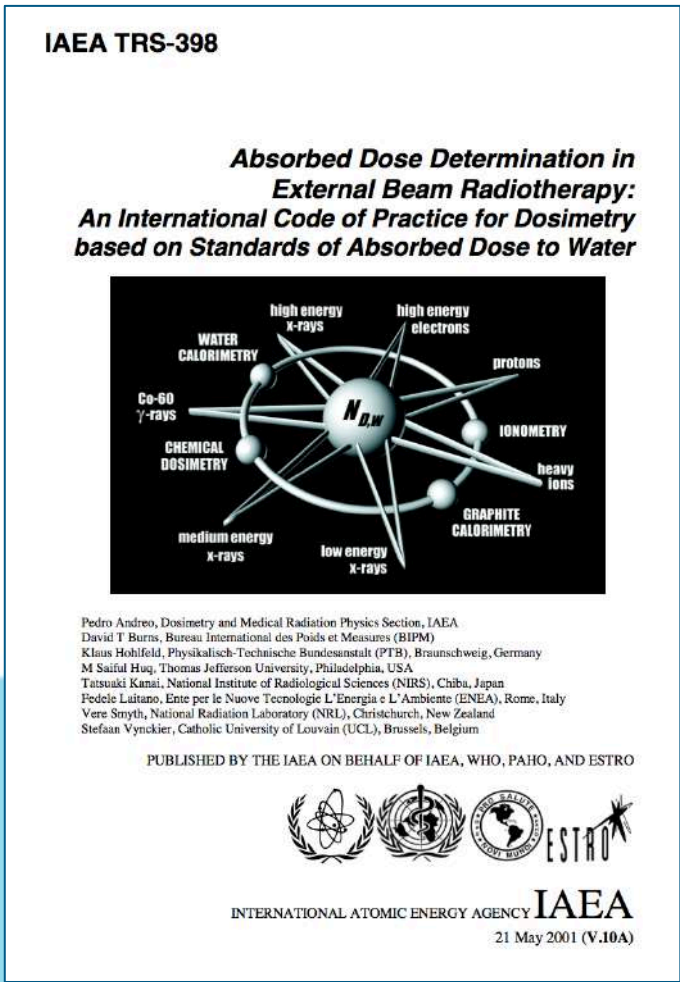
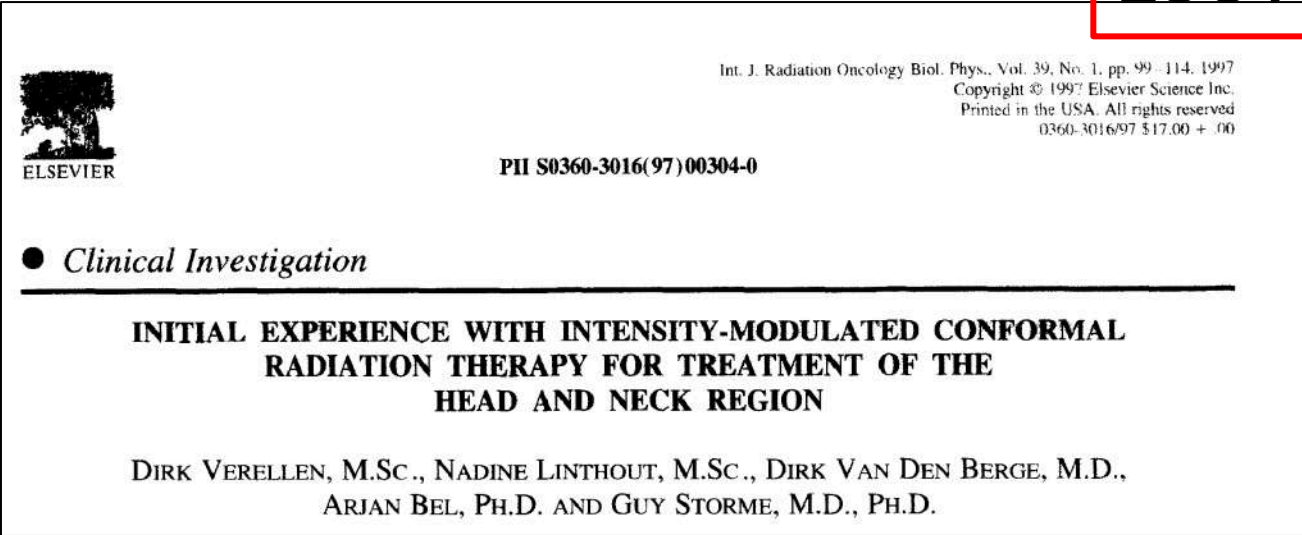
Das IJ, et al., Med Phys, 2008

D. Verellen - What if no standards?



Absolute dose calibration

1997

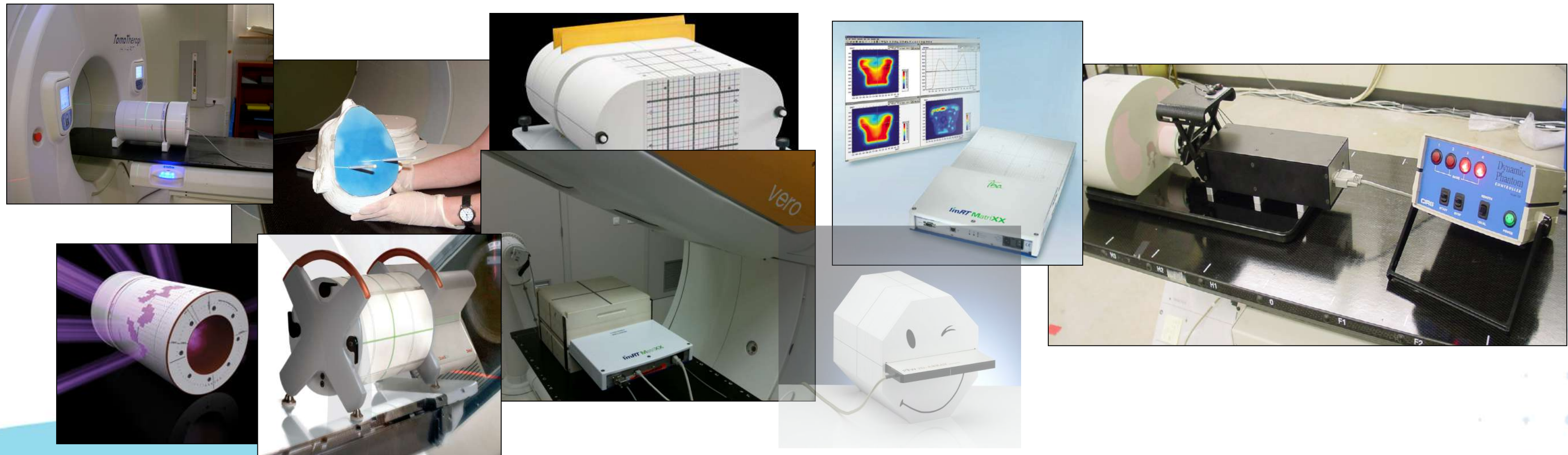


2021

2017

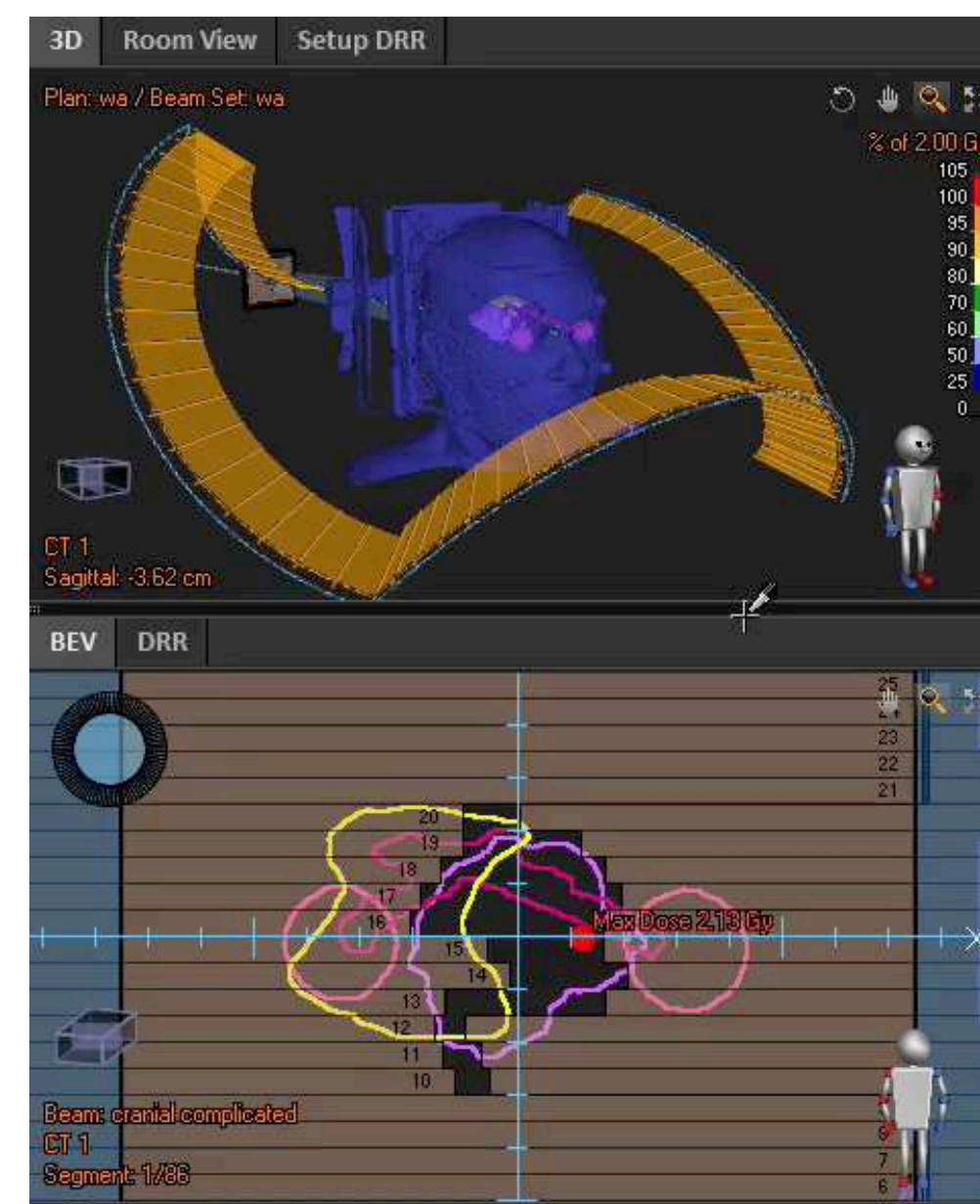
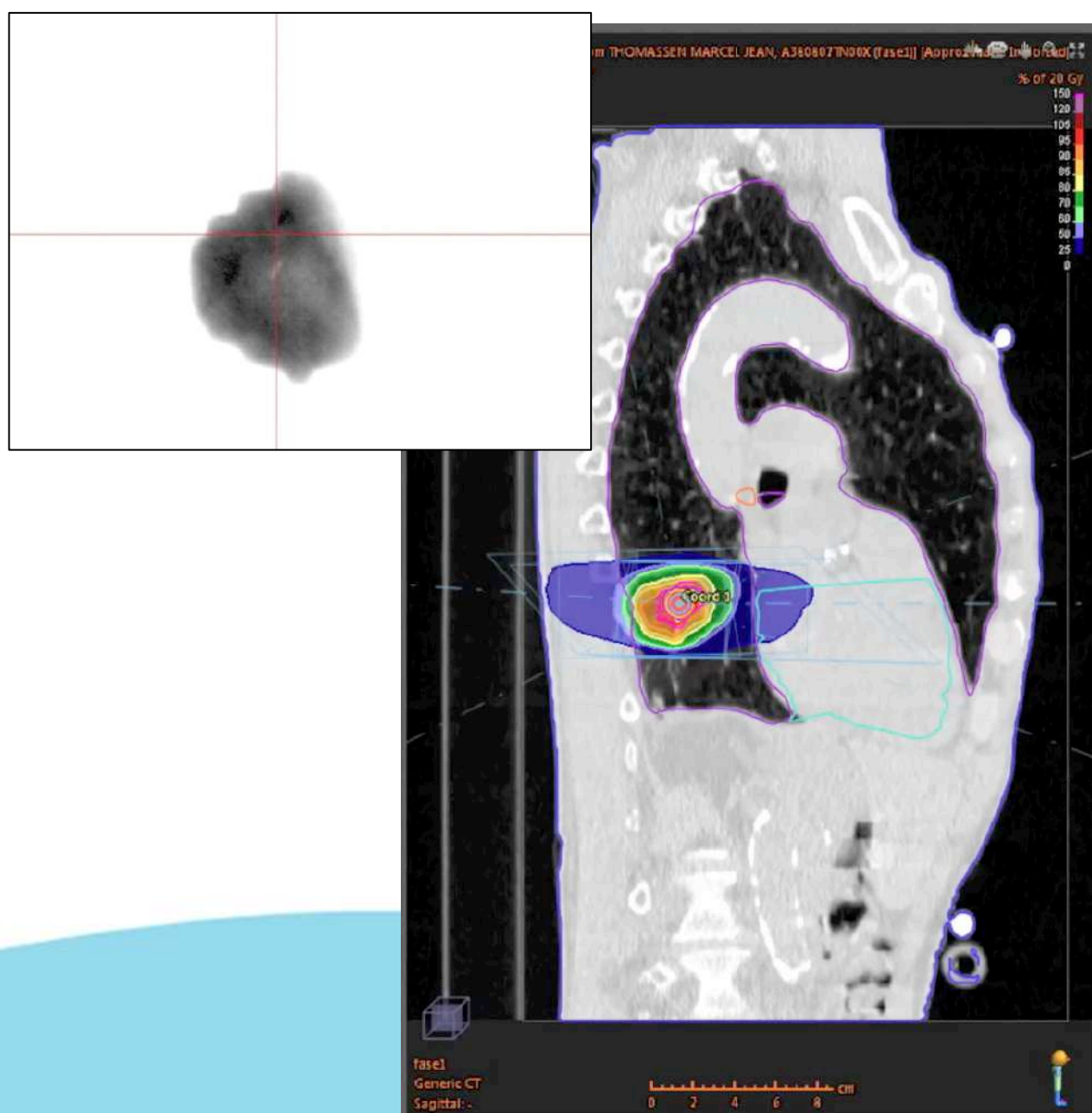
Patient specific QA

In vivo dosimetry was no longer feasible ... so we “exaggerate” commissioning ...



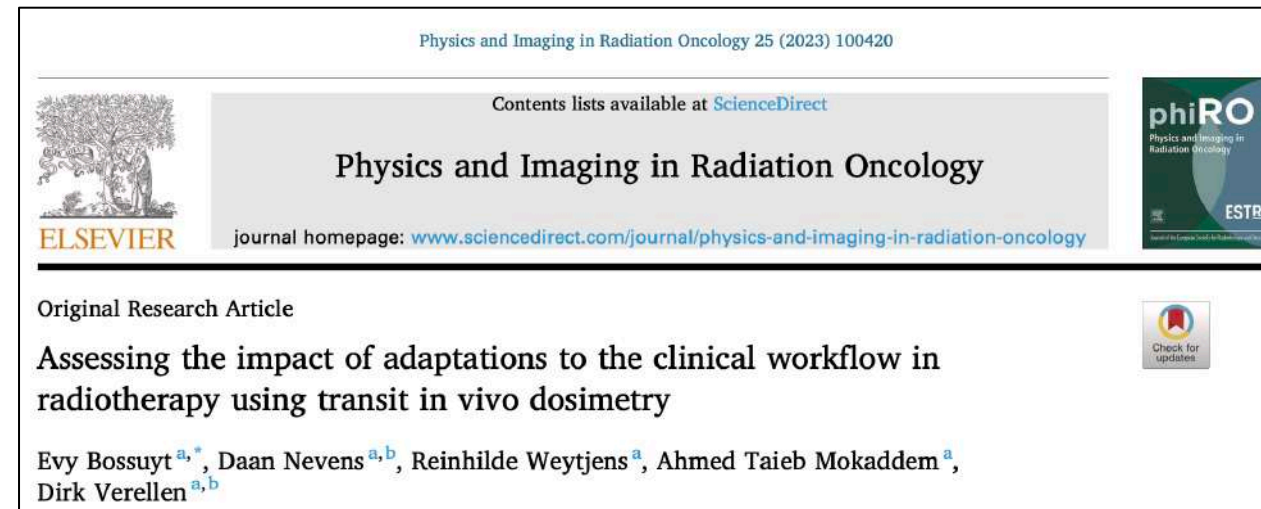
Patient specific QA

What about (manual) independent MU verification?

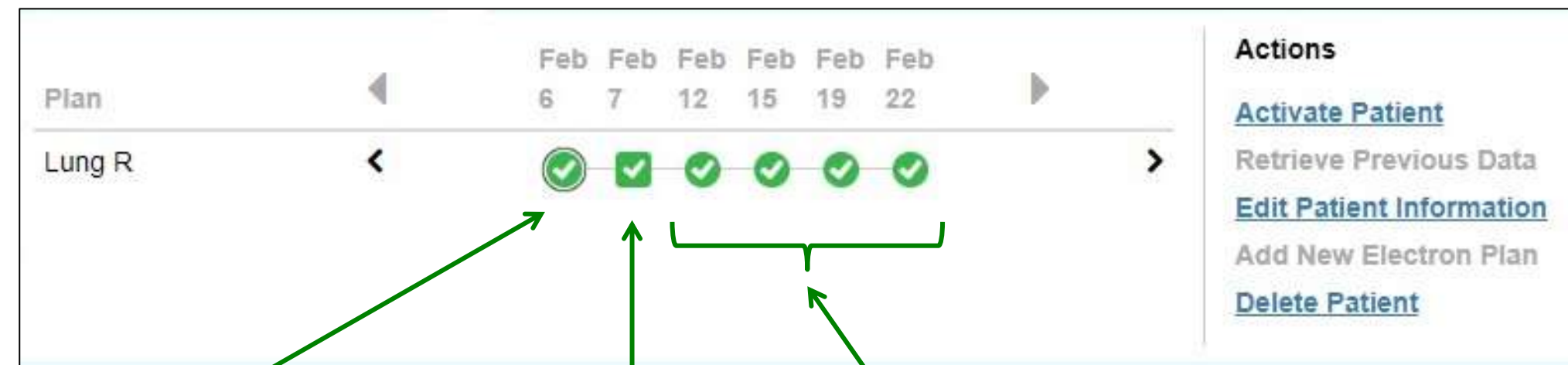


Patient specific QA

Patient specific QA ... Today



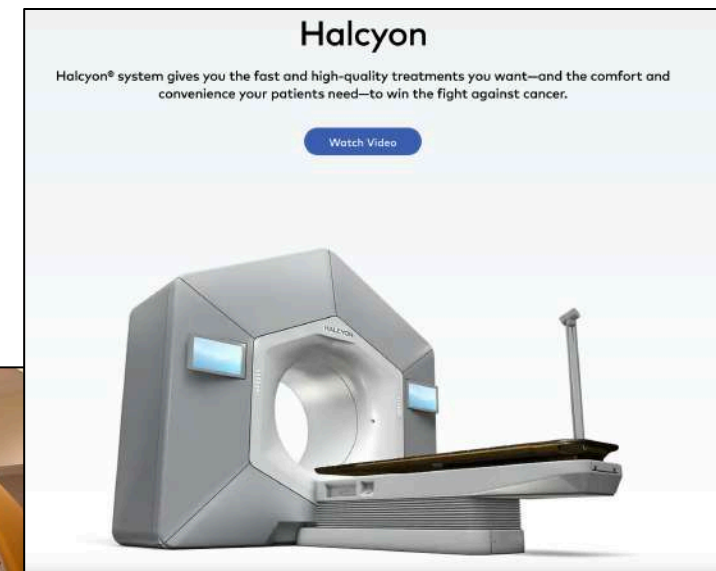
- Patient-specific pre-treatment QA and *in vivo* dosimetry are performed for every patient using the PerFraction platform (Sun Nuclear), which includes:



- Independent MU calculation,
- Dose transmission measurements, pre-treatment,
- Dose transmission measurements, during treatment (*in vivo*)

Implementing a new “established” technique

- Vendors are selling complex equipment as **plug&play** to facilities with **understaffed-undertrained** staff.
- Hospitals are using internal **procedures and workflows** that are **not adapted** to these new technologies.
- Software and hardware is continuously being **upgraded**.



Radiotherapy Workflow



Radiotherapy Workflow ... all in one?



Efficiency, Standardization, Quality & Safety



AORN JOURNAL THE OFFICIAL VOICE OF PERIOPERATIVE NURSING

Patient Safety First

Using Aviation Safety Measures to Enhance Patient Outcomes

Russell M. Rivers, Diane Swain RN, William R. Nixon

First published: 01 January 2003 | [https://doi.org/10.1016/S0001-2092\(06\)61385-9](https://doi.org/10.1016/S0001-2092(06)61385-9) | Citations: 18

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ABSTRACT

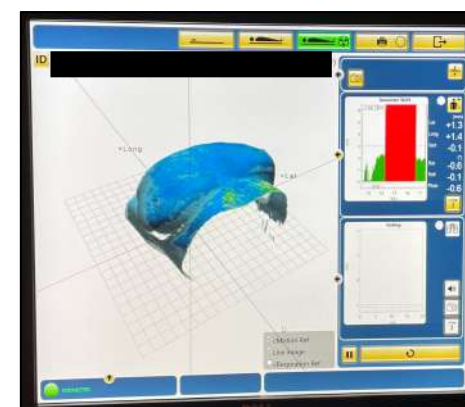
Recent media reports have put a spotlight on the increasing number of medical errors occurring in US health care institutions. In contrast to health care's increasing error rate, the aviation industry is experiencing a decreasing error rate. Could the safety techniques used in the aviation industry be applied to health care? This article explores that question. The dynamics of the surgical suite are not unlike those of the cockpit of an airplane; therefore, perioperative services was selected to pilot test the aviation model of safety training. *AORN J* 77 (Jan 2003) 158-162.

Automate or perish

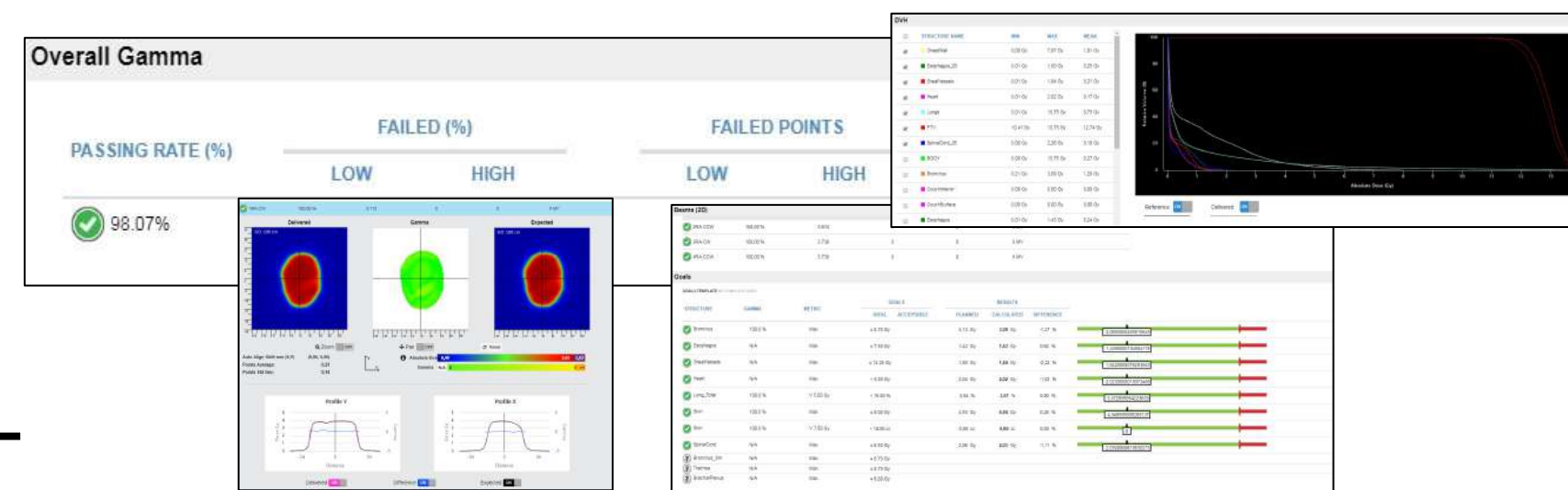
Standardization
&
Automation



Efficiency & Optimization

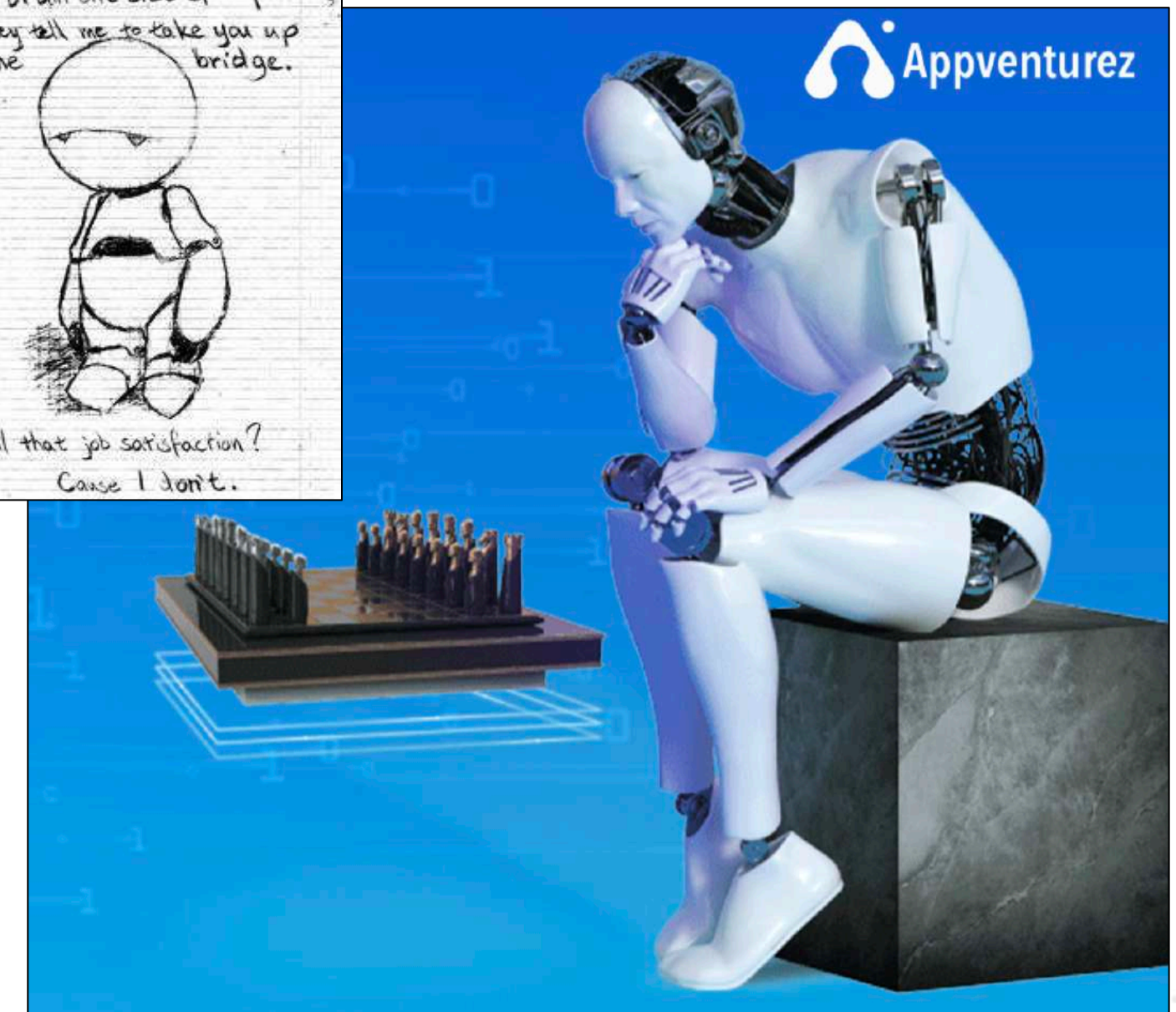
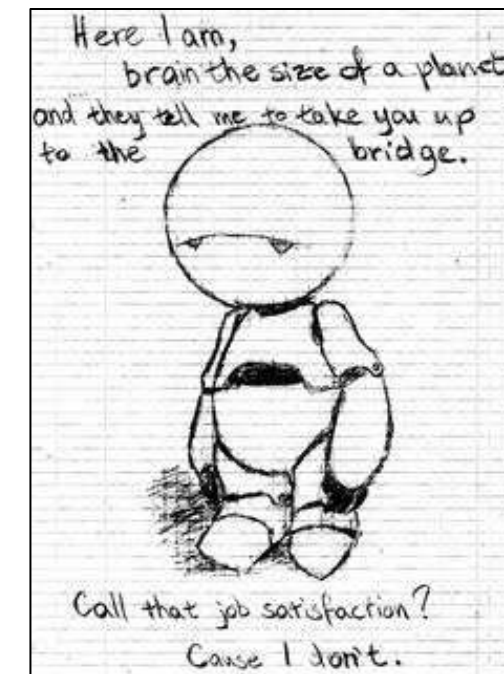


Safety & Quality



Complexity increases ... enters automation ...

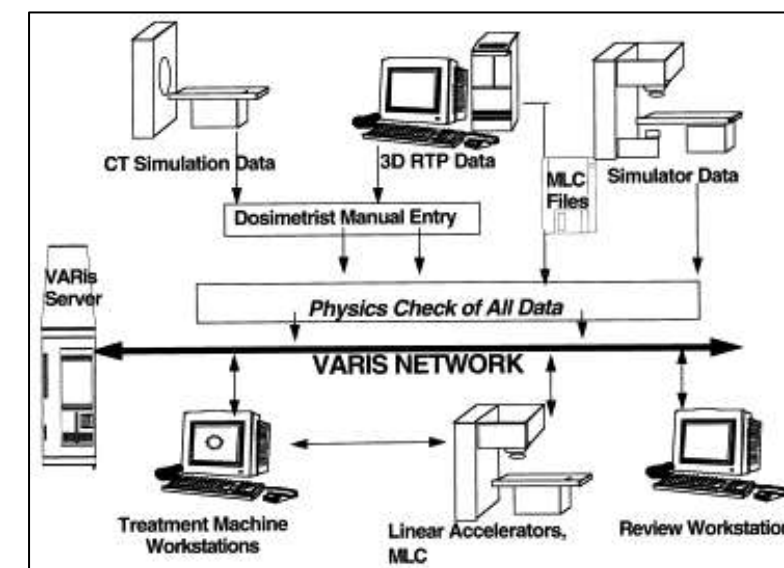
**With large scale introduction of
scripting and AI
(computerized black boxes),
the role of human interaction needs
rethinking**



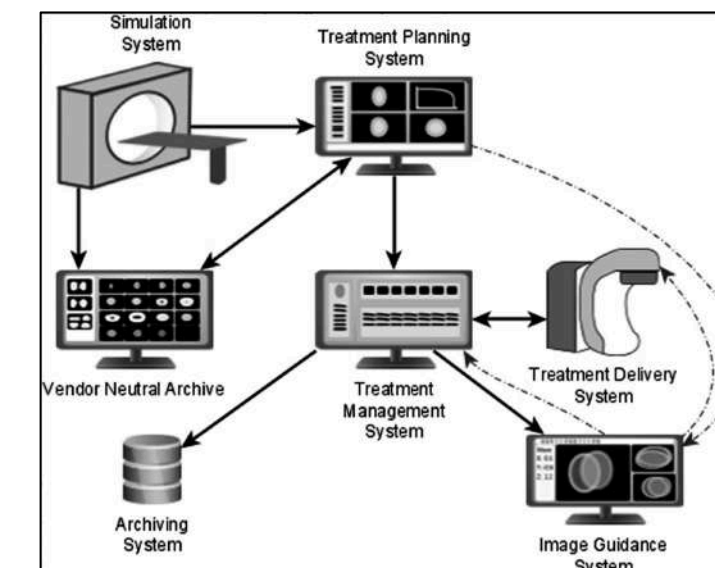
Complexity increases ... enters automation ...

Similar to the introduction of R&V systems at the end of the previous century:

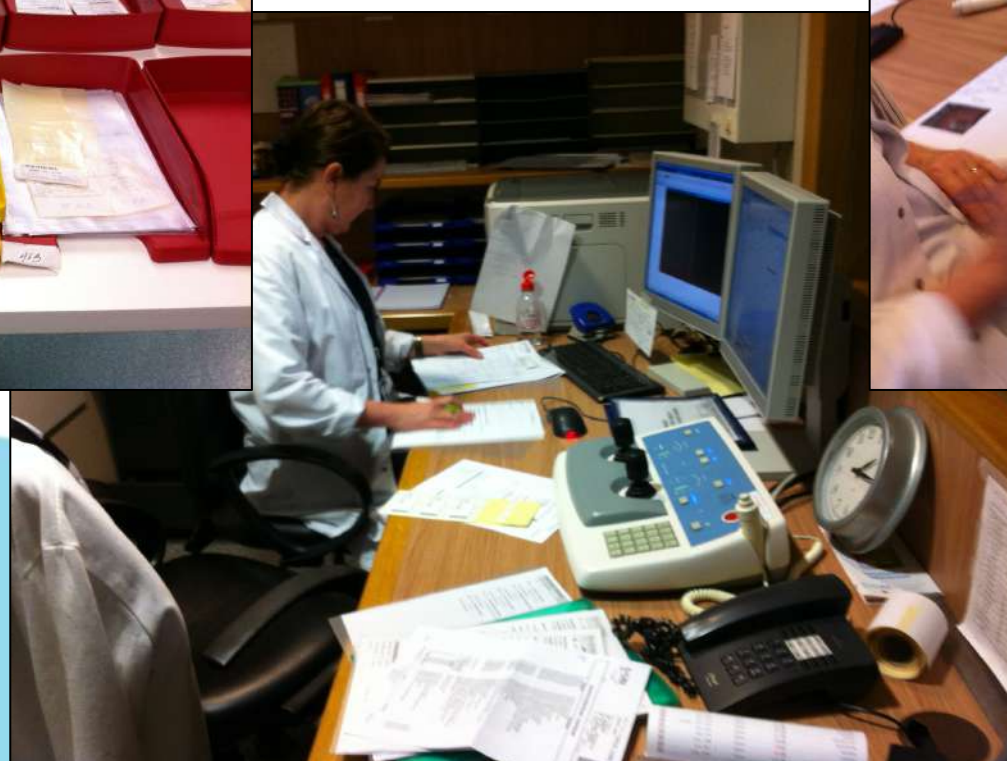
- It was introduced to *mitigate human errors and reduce repetitive tasks*
- But people started to verify these tasks as if it was human, *... by performing repetitive tasks ...*



IJROBP



AAPM TG201



Standards on MSQA and PSQA?

Hypothetical situation ...

What if during an audit, a centre claims that **MSQA can be minimized**?

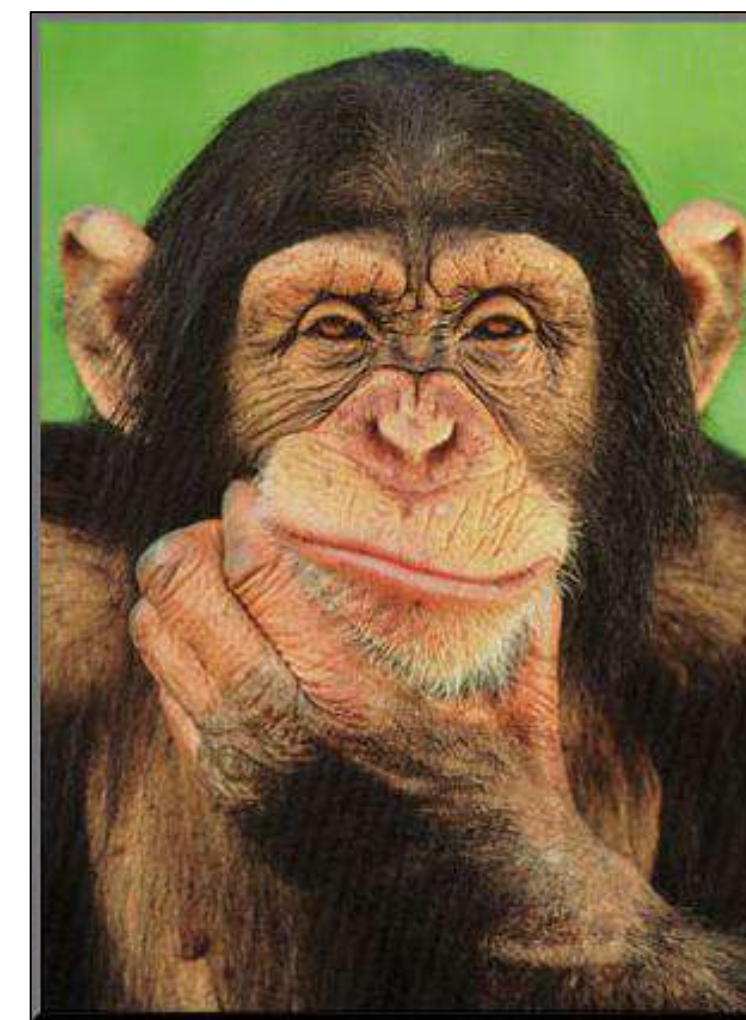
- Modern technology comes with efficient self-tests, the system will have an interlock before our MSQA detects it,
- Service engineers “know” what they’re doing,

What if the centre also claims, **independent dose calculation is useless**, because:

- The algorithm is less accurate than the clinical algorithm,
- Redundancy after upgrades is tested, so the accuracy is maintained,

What if the centre **omits PSQA and IVD** if the plan fits criteria of class solution?

- PSQA is only re-commissioning,
- EPID-IVD is not sensitive-specific, with many falls-positives / falls-negatives.



Standards on MSQA and PSQA?

- **MSQA:**
 - With increasing techniques and technology, we add more and verification tests ... **redundancy versus efficiency**
 - Vendors and equipment become more reliable and come with efficient “**self-tests**” ... user specific MSQA almost never discovers “new” errors.
 - **How to maintain “independent” QA?**
- **PSQA:**
 - Pre-treatment QA does not include the patient, hence is it not merely **re-checking commissioning**?
 - With **intelligent complexity metrics, class solutions and AI**: can't we abandon or at least reduce PSQA?
 - 99.99% of PSQA pass ... **it's a waste of time?**
 - **IVD is really PSQA as it includes the patient**: but if an error occurs, it's too late (especially as we tend to hypofractionate or introduce oART ...)
- Re-evaluate the combination MSQA-PSQA on a regular basis:
 - By default standards are obsolete?
 - Ask yourself: **if an error occurs, will the MSQA-PSQA catch it?** ... FMEA!!!



Franquin

What are we lookin' for?



Ensuring that *what has been planned, really has been delivered!!*

- ... *for each patient!!*
- ... *every fraction!!*
- ... *in a safe way!!*

Due to the complexity of Radiation Oncology, QA and safety is more than pre-treatment verifications, but process oriented.

Dosimetry audits and standards ...

- “An **external dosimetry audit** is mandatory **PRIOR** to clinical implementation of a linac”

- What are the tolerances, who decides on these tolerances?
- Basic dosimetry audit or complex techniques?
- What if audit fails for, let's say 1 electron energy only?
- What in case of dispute, experimental set-up errors?



See presentation on “How to maintain a permanent audit service”

Summary

- The audit is NOT a pass/fail exam, it's advisory
- Focus on patient safety and quality
- It's more than MSQA and PSQA: process oriented QA





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