Commentary

On the Need for Better Informing Results From Randomized Clinical Trials in Oncology

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Cancer patients frequently seek information from their oncologists regarding the anticipated outcomes of their treatment. They commonly inquire about the potential for a cure and the likelihood of achieving it, as well as whether the treatment will prolong their life and, if so, by how much. In oncology, randomized controlled trials play a pivotal role in informing treatment decisions. It is essential that published randomized controlled trials provide comprehensive data to enable clinicians to deliver precise and reliable information to patients. The concept of the Hazard Ratio (HR) can be challenging for many oncologists and is often misunderstood in terms of its implications for risk. Thus, incorporating both relative and absolute risk metrics in addition to hazard ratios could substantially enhance patient care. This consideration is crucial not only for patients and oncologists but also for policymakers within public health systems.

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Introduction

Randomized controlled trials (RCTs) are currently the most widely utilized and regarded as the gold standard for providing robust evidence in clinical decision-making, although they are not immune to biases that may influence therapeutic recommendations. Given the often-fatal nature of cancer, patients seek the clearest and most definitive information about the impact of prescribed treatments on their disease prognosis. Common inquiries from patients include: "Can this treatment cure me, and what is the likelihood of that?" and "Will this treatment extend my life, and if so, by how much?" From a public

health perspective, particularly in oncology where treatments are often costly, it is essential for decision-makers to have comprehensive data on the effects of oncological interventions on survival outcomes.

To assess how RCT results in oncology are presented in the academic literature, we examined the most recent five RCTs involving malignant solid tumors published in the New England Journal of Medicine from June to August 2024. The selection was done by scanning the NEJM content from the more recent date (August 2024) backward (June 2024). The five most recent randomized studies on solid tumors published were selected without any additional consideration.

Published RCT in the NEJM

1. The first study compared Beltizufan and Everolimus in patients with advanced renal cell carcinoma[1]. This study enrolled 374 patients receiving Beltizufan and 272 receiving Everolimus. The dual primary endpoints were progression-free survival (PFS) and overall survival (OS). In the first interim analysis, at a median follow-up time of 18.4 months, the median PFS was 5.6 in both arms (95% CI 3.9-7 and 4.8-5.8 respectively) and a HR of 0.75 (95% CI 0.63-90), however, curves crossed, breaking the proportional hazards assumption. For OS, at a median follow-up of 25.7 months, the median was 21.4 months in the Beltizufan group and 18.1 months in the Everolimus group (HR for death, 0.88; 95% CI, 0.73-1.07; p=0.20).

The number of PFS events was 289/374 and 276/372 respectively (Figure S3A, suppl data, appendix). These numbers translate into a Relative Risk (RR) for progression of 1.041% (95% CI 0.960 - 1.129), RR increase of 4.10%, p=0.3270, and an increase of Absolute Risk (AR) for progression of 3.08% (95% CI, -3.07-9.23). Accordingly, the number needed to harm (NTH) is 32. Regarding the RR and AR for death (213 and 228 deaths in the Beltizufan and Everolimus groups respectively, data in text), the numbers are RR 0.923 (95% CI 0.824- 1.047), a reduction of 7.81%, p=0.2286, while the AR reduction is 4.34% (-3.07%-9.23%), both lacking statistical significance. The number needed-to-treat (NNT) is 23 (95% CI, 8.1-Infinity).

2. This study evaluated the efficacy of adjuvant Osimertinib versus placebo following chemoradiation in patients with Stage III EGFR-mutated non-small cell lung cancer[2]. Of the 143 participants, 73 were assigned to the placebo group, while the remaining 70 received Osimertinib. The results showed a significant PFS benefit with Osimertinib, as the median PFS was 39.1 months compared to 5.6 months with placebo. The HR for disease progression or death was 0.16 (95% CI, 0.10-0.24; p<0.001). Interim OS

data revealed a 36-month OS among 84% of patients with Osimertinib and 74% with placebo, with a HR for death of 0.81 (95% CI, 0.42-1.56; p=0.53).

The study reported 57 PFS events out of 143 in the osimertinib group and 63 out of 73 in the placebo group (text results). These results correspond to an RR for progression of 0.461% (95% CI: 0.3703- 0.5762; p<0.0001) and an AR reduction of 46.40% (95% CI: 35.19-57.69%). This indicates a high efficacy of Osimertinib, with an NNT of 2, meaning that treating two patients with Osimertinib is required to prevent one progression event.

However, the impact on mortality is less pronounced. There were 28 and 15 deaths in the osimertinib and placebo groups, respectively (Overall Survival Suppl results, appendix). The RR for death is 0.952% (95% CI, 0.5442-1.6686; p=0.866), which is not statistically significant (RR reduction 4.80%), while the AR reduction is 3% (95% CI, -10.36-12.29%). Consequently, the NNT to prevent one death is 104 (95% CI, 8.1-Infinity), indicating that 104 patients would need to be treated with Osimertinib to avoid one death.

3. In the study of Tisotumab Vedotin[3], a total of 502 patients were randomized, with 253 assigned to receive Tisotumab Vedotin and 249 assigned to receive chemotherapy for recurrent cervical cancer. The primary endpoint was OS. At a median follow-up time of 10.8 months, the median OS was significantly longer in the Tisotumab Vedotin group than in the chemotherapy group (11.5 months 95%CI, 9.8 to 14.9 vs. 9.5 months 95% CI, 7.9-10.7), results that represented a 30% lower risk of death with Tisotumab Vedotin than with chemotherapy (HR, 0.70; 95% CI, 0.54- 0.89; p=0.004).

As shown in the PFS curve, the number of events was 198/253 and 194/249 for the Tisotumab Vedotin and chemotherapy groups, respectively (Figure 3A). This translates into an RR of 1.004 (95%CI 0.915 - 1.102), p = 0.095), RR increase of 0.004% while AR increase was 0.35% % (95% CI -6.89 -7.59) for a NNH of 286 (95% CI 13.2 -Infinity). Neither the RR nor the AR reductions for PFS are statistically significant. Regarding OS, there were 123/253 and 140/249 events of death for Tisotumab Vedotin and chemotherapy, respectively (Figure 2B). With these numbers, the RR for death is 0.8647 (9%CI 0.7313 to 1.0224), p=0.0889, with a RRr of 13.5%, while the AR reduction is 7.61% (-1.10 − 16.32) and NNT 12 (6.1-Infinity). None of these are statistically significant.

4. In this RCT[4] involving patients with previously untreated EGFR-mutated advanced NSCLC, 429 patients were randomized to the Amivantamab-Lazertinib group, 429 to the Osimertinib group, and 216 to the Lazertinib group. The median PFS was significantly longer in the Amivantamab−Lazertinib group. The primary endpoint was PFS in the Amivantamab−Lazertinib group compared to Osimertinib. At a

median follow-up time of 22 months, the median PFS was significantly longer in the Amivantamab–Lazertinib group than in the Osimertinib group (23.7 vs. 16.6 months); HR for disease progression or death, 0.70; 95% CI, 0.58 – 0.85; p<0.001). In the planned interim analysis, the HR for death with the combination was 0.80 (95% CI, 0.61 – 1.05).

The number of PFS events was 192/429 in the Amivantamab-Lazertinib group and 252/429 in the Osimertinib group (Figure 1C). These numbers translate into an RR for progression of 0.761 (95% CI 0.6679 – 0.8691), p=0.0001, for a RRr of 23.90%. The AR reduction for progression was 5.24% (95% CI –1.17 – 11.66%). This corresponds to an NNT of 20 (95% CI, 8,6 – Infinity).

Regarding OS, there were 97 and 117 deaths for Amivantamab-Lazertinib and Osimertinib, respectively (data in the text). The RR for death was 0.829 (95%CI 0.656 –1.047), p =0.116, for a RRr of 17.1%, while for AR reduction it was 4.66% (95% CI, –1.12 –10.44), respectively, with an NNT of 22 (95% CI 9.6 – Infinity), none of which were statistically significant.

5 In this randomized controlled trial (RCT) evaluating adjuvant therapy in resected stage III melanoma patients harboring BRAFV600 mutations[5], 438 patients were allocated to the Dabrafenib plus Trametinib arm, while 432 patients were assigned to the placebo arm.

At a median follow-up of almost ten years, the estimated OS at 8 years was 71% with Dabrafenib plus Trametinib and 65% with placebo (HR for death, 0.80; 95% CI, 0.62 – 1.01; p=0.06). PFS also favored Dabrafenib plus Trametinib, 93.1 vs. 16.6 months, HR 0.52 (95% CI, 0.43-0.63). According to the number of events of PFS (438/198 and 432/264 respectively), Figure 3A, the RR was 0.739 (95%CI, 0.651-0.849), p<0.0001, for a RRr of whereas the AR reduction was 15.91% (95%CI, 9.36-22.45) for a NNT of 7 (95%CI, 4.5-10.7). Regarding OS, the number of events were 125/438 and 136/432 for Dabrafenib plus Trametinib and placebo, respectively (Figure 1A). The RR was 0.906 (95% CI 0.7398 1.1109), p=0.344, for a RRr of 9.40%, whereas the AR reduction was 2.94% (95% CI –3.15 – 9.03) and a NTT of 34 (95% CI, 11.1 to Infinity). Both risk reductions had no statistical significance.

Comments

The publication of RCT results in cancer treatment through scientific journals and mainstream media plays a pivotal role in empowering both oncologists and patients. It provides the necessary information to address two fundamental patient questions: "Doctor, can this treatment cure me, and what is the

likelihood?" and "Doctor, can this treatment extend my life, and by how much?" Accurate and thorough reporting of these outcomes is essential for informed decision-making in clinical practice.

Most oncologists are familiar with measures such as median survival and 1- or 5-year survival estimates, typically obtained using the Kaplan and Meier method. A simple initial analysis of survival usually involves calculating and graphically presenting the Kaplan-Meier survival estimates for each treatment regimen. In addition, estimates of the median survival (with confidence intervals) or 1-, 3-, or 5-year survival probabilities (with standard errors) can be presented, followed by the use of the log-rank test to assess the significance of any differences observed in these survival curves. Then, the Cox proportional hazards regression method assesses treatment outcomes after adjustment for multiple covariates. As a result, HRs are commonly used to estimate the treatment effect in oncology RCTs.

Despite the common use of HRs in reporting results from oncology RCTs, they remain one of the most perplexing concepts for clinicians. The HR estimates the hazard rate ratio between the experimental and control groups over the entire study duration. The ratio is estimated at different time points along the entire time of the curve, and although the ratio can change over time, a constant HR is calculated. A simplistic interpretation is that an HR of 1 means equal efficacy between the experimental and control treatments; an HR lower than 1 indicates that the experimental arm is better, and an HR higher than 1 indicates that it is worse than the control. It is important to note that HR is a relative measure. In a hypothetical case where an RCT would not show the survival curve, nor mention the median OS and OS percentages at different time points, it is impossible to determine the magnitude of the survival effect of treatment if only the HR is provided. Nevertheless, there is a direct relationship between the magnitude of HR reduction and increased survival, but this very much depends on the survival of the control arm. For instance, a HR of 0.20 translates into a high increase in survival if the control arm has a 2-year survival rate of 10%, but a much lower increase if the control arm has a 2-year survival rate of 90%.

In other words, estimating the magnitude and direction of survival outcomes in RCT requires information on 1) the HR, 2) the median survival time, and 3) the percentage of patients alive at different time points, i.e., 1-year, 3-year, and 5-year. These concepts are nicely explained in[6][7][8]. While points 2 and 3 are straightforward in their meaning, special care is needed to interpret the common statements on HR with published RCT, which can be as follows: Drug X showed an HR of 0.75, which reduced the risk of death by 25%. While the statement is correct, this reduction in the risk of death should not be interpreted as a risk reduction in the commonly understood sense. If not further explained, the statement conveys an implied durability of the effect, which can be erroneously interpreted to believe that Drug X can eliminate

the chance of death for a fraction of patients. Indeed, the reduction of the HR means that survival is prolonged but not that the risk of death in such a proportion has been averted; in simpler terms, the Drug X HR reduction means a reduction in the rate of events that happen. The risk of death, as commonly understood, must be calculated based on the actual number of patients alive or dead at a fixed time point (i.e., when the study is closed or at the time it is reported). The RR and the AR must be determined, the first based on percentages and the second in absolute numbers. Unfortunately, in this sample of 5 RCT, only two (Tisotumab Vedotin and Dabrafenib/tTametinib) studies disclosed the number of events in the curve figures, which makes it difficult for readers to calculate Relative and Absolute risks, despite these metrics being a must for RCT according to the CONSORT statements[9].

As demonstrated in **Table 1**, while all five studies reported a statistically significant reduction in the hazard ratio (HR) for progression-free survival (PFS), only 3 (osimertinib, amivantamab, and dabrafenib studies) statistically significantly reduced the RR, whereas in the remaining two studies, the RR actually increased, though non-statistically significantly. Likewise, the PFS Absolute Risk was only statistically significantly reduced in the osimertinib and dabrafenib studies. Regarding the risk of death, the HR was only statistically significantly reduced in the tisotumab trial, but in none of the five were either the Relative or Absolute risk for death reduced.

If the publication of RCTs routinely included comprehensive information on 1) HR, 2) median OS, and 3) OS proportions at various time points (e.g., 6 months, 1 year, 3 years, 5 years), along with the associated RR, AR, and NNT derived from AR reduction, oncologists would be better equipped to address patients' questions regarding the expected outcomes of cancer treatments. Furthermore, given the increasing dissemination of RCT results through mainstream media, presenting these findings in such a thorough manner would also enhance patients' understanding of their treatment options.

The authors of this work, though not expert statisticians, have endeavored to elucidate these complex concepts. However, it is possible that some nuanced details and potential pitfalls in the calculation and application of hazard ratios, relative risk, absolute risk, and number needed to treat in time-to-event analysis may have been overlooked[10][11]. Nonetheless, it is imperative that the legitimate interests of the pharmaceutical industry and journal editors do not supersede the primary goal of advancing patient care in oncology. Your engagement in this discussion is vital to ensuring that the interests of cancer patients remain at the forefront.

Study	PFS HR	PFS RRr%	PFS ARr%	OS HR	OS RRr%	OS ARr%	OS NNT
Beltizufan	0.75*	+4.10	+3.08	0.88	7.81	4.34	23
Osimertinib	0.16*	53.89*	46.40*	0.81	4.80	0.97	104
Tisotumab Vedotin	0.67*	+0.004	+0.35	0.70*	13.5	7.61	14
Amivantamab/Lazertinib	0.70*	23.90*	5.24	0.80	17.1	4.66	22
Dabrafenib/Trametinib	0.52*	26.1*	15.91*	0.80	9.40	2.94	34

Table 1. Hazard Ratio, Relative Risk, and Absolute Risk reductions and NNT for survival.

*Numbers and percentages are shown without 95% CI for better appreciation. Bold and *: statistically significant.*

Numbers with + mean that the RR actually increased; none of these increases are statistically significant.

Readers can easily calculate them using free online calculators.
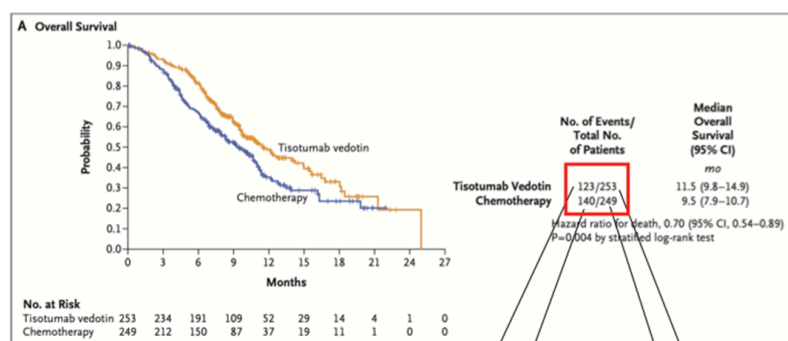
Relative Risk Online Calculator

https://www.medcalc.org/calc/relative_risk.php

Absolute Risk Online Calculator

http://araw.mede.uic.edu/cgi-bin/nntcalc.pl

Additional Information

Figure demonstrating how to calculate the Absolute Risk. Data taken from the Tisotumab Vedotin trial.

A Overall Survival

	No. of Events/ Total No. of Patients	Median Overall Survival (95% CI)
		mo
Tisotumab Vedotin	123/253	11.5 (9.8–14.9)
Chemotherapy	140/249	9.5 (7.9–10.7)

Hazard ratio for death, 0.70 (95% CI, 0.54–0.89)
P=0.004 by stratified log-rank test

No. at Risk

	0	3	6	9	12	15	18	21	24	27
Tisotumab vedotin	253	234	191	109	52	29	14	4	1	0
Chemotherapy	249	212	150	87	37	19	11	1	0	0

Total - events 253 - 123 = **130** 249 - 140 = **109**	Good Outcome **Alive**	Bad Outcome **Dead**	Lost to follow-up (optional)	Total
New Therapy	130	123		253
Control	109	140		249
Total	239	263	0	502

If the curve does not show the number of events, it can be found in the PFS or OS forest plot in prespecified subgroups. If it is not there, the number of events can be described in the text's results. If not there, it can be in the "appendix" or supplementary information. If not there, then it is not showed.

Absolute risk reduction: **7.61%** of patients will not experience adverse events (**death**) under New Therapy that they would have under Control.

95% CI: [-1.10%, 16.32%]

Note that the confidence interval crosses zero, which implies that the absolute risk reduction is not significant. New Therapy may actually increase risk.

Number needed-to-treat (NNT): You must treat **13.1** patients with New Therapy to prevent 1 adverse event (**death**) that would have happened under Control. To be conservative, you may choose to round this number **up** to the nearest patient. 95% CI: [6.1, Infinity]

("Risk" as used above refers to the chance of experiencing a adverse event.)

This webpage of the BMJ Best Practice is recommended for help understanding statistical risks.

https://bestpractice.bmj.com/info/us/toolkit/learn–ebm/how–to–calculate–risk/

References 10, 11, and our previous work (*Alfonso Dueñas-Gonzalez. (2024). Risk of death derived from Hazard Ratio should not be communicated as Relative Risk reductions for death in cancer clinical trials. Intentional or inadvertent?. Qeios. doi:10.32388/F3BC9A.3*) can be of help for better understanding the meaning of risk in clinical trials.

Statements and Declarations

Data and Software Availability

No data are associated with this article.

Competing Interests

The author declares not to have conflicts of interest.

Author Contribution

The content and ideas expressed in this work are the sole responsibility of the authors. Both authors contributed equally to this work.

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Declarations