



DIGITAL ACCESS TO SCHOLARSHIP AT HARVARD

Why tyrosine kinase inhibitor resistance is common in advanced gastrointestinal stromal tumors

The Harvard community has made this article openly available.
[Please share](#) how this access benefits you. Your story matters.

Citation	Tomasetti, Cristian, George D Demetri, and Giovanni Parmigiani. 2013. "Why tyrosine kinase inhibitor resistance is common in advanced gastrointestinal stromal tumors." F1000Research 2 (1): 152. doi:10.12688/f1000research.2-152.v1. http://dx.doi.org/10.12688/f1000research.2-152.v1 .
Published Version	doi:10.12688/f1000research.2-152.v1
Accessed	April 17, 2018 4:42:06 PM EDT
Citable Link	http://nrs.harvard.edu/urn-3:HUL.InstRepos:11879540
Terms of Use	This article was downloaded from Harvard University's DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA

(Article begins on next page)



SHORT RESEARCH ARTICLE

Why tyrosine kinase inhibitor resistance is common in advanced gastrointestinal stromal tumors [v1; ref status: indexed, <http://f1000r.es/1ac>]

Cristian Tomasetti^{1,2,5,6}, George D Demetri^{3,4}, Giovanni Parmigiani^{1,2}

¹Department of Biostatistics, Harvard School of Public Health, Boston MA, 02215, USA

²Department of Biostatistics and Computational Biology, Dana-Farber Cancer Institute, Boston MA, 02215, USA

³Ludwig Center for Cancer Research at Dana-Farber Cancer Institute and Harvard Medical School, Boston MA, 02215, USA

⁴Center for Sarcoma and Bone Oncology, Dana-Farber Cancer Institute, Boston MA, 02215, USA

⁵Current affiliation: Division of Biostatistics & Bioinformatics, Sidney Kimmel Comprehensive Cancer Center, Johns Hopkins University School of Medicine, Baltimore, MD , 21205-2013 , USA

⁶Current affiliation: Department of Biostatistics, Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, 21205-2179 , USA

v1 **First Published:** 09 Jul 2013, 2:152 (doi: 10.12688/f1000research.2-152.v1)
Latest Published: 09 Jul 2013, 2:152 (doi: 10.12688/f1000research.2-152.v1)

Abstract

Background: Most patients with advanced gastrointestinal stromal tumors (GIST) develop drug resistance to tyrosine kinase inhibitors (TKIs) within two years of starting therapy, whereas most chronic myeloid leukemia (CML) patients in chronic phase still exhibit disease control after a decade on therapy. This article aims to explain this divergence in long term outcomes.

Methods and results: By combining clinical and experimental observations with mathematical formulas we estimate that, in advanced GIST, the genetic changes responsible for resistance are generally already present at disease detection.

Conclusion: This result has relevant clinical implications by providing support for the exploration of combination therapies.

Article Status Summary

Referee Responses

Referees	1	2
v1 published 09 Jul 2013	 report 1	 report

1 **Shiro Urayama**, University of California Davis Medical Center USA

2 **Gerald Prager**, University of Vienna Austria

Latest Comments

No Comments Yet

Corresponding author: Cristian Tomasetti (ctomase2@jhmi.edu)

How to cite this article: Tomasetti C, Demetri GD, Parmigiani G (2013) Why tyrosine kinase inhibitor resistance is common in advanced gastrointestinal stromal tumors [v1; ref status: indexed, <http://f1000r.es/1ac>] *F1000Research* 2013, 2:152 (doi: 10.12688/f1000research.2-152.v1)

Copyright: © 2013 Tomasetti C et al. This is an open access article distributed under the terms of the [Creative Commons Attribution Licence](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited. Data associated with the article are available under the terms of the [Creative Commons Zero "No rights reserved" data waiver](#) (CC0 1.0 Public domain dedication).

Grant information: The work of CT was supported in part by the National Institute of Health under Grant T32 CA009337. GDD was partially funded by DFHCC SPORE (NCI Grant) for GIST research, as well as Ludwig Center for Cancer Research at Dana-Farber/Harvard Cancer Center. GP was supported in part by the NIH/NCI grant 5P30 CA006516-46.

The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests:

CT and GP have no competing interests to disclose. GDD discloses the following: Novartis. Consultant, <\$10k per annum. Honorarium <\$10k per annum. Research support to Dana Farber for specific clinical trial agreements in our sarcoma unit, >\$200,000 per year. Research support to Dana-Farber for specific clinical trial agreements in our sarcoma unit, >\$200,000 per year. Pfizer. Consultant, <\$10k per annum. Honorarium <\$10k per annum. Research support to Dana-Farber for specific clinical trial agreements in our sarcoma unit, >\$200,000 per year. Infinity Pharmaceuticals. Consultant, <\$10k per annum. Research support to Dana-Farber for specific clinical trial agreements in our sarcoma unit, >\$200,000 per year. Regulatory presentation support, uncompensated. Glaxo Smith Kline. Consultant, <\$10k per annum. Kolltan Pharmaceuticals. Chair, Scientific Advisory Board. Chair, Medical Advisory Board. Consultant. Equity (minor stake, non-public).

First Published: 09 Jul 2013, 2:152 (doi: 10.12688/f1000research.2-152.v1)

First Indexed: 18 Jul 2013, 2:152 (doi: 10.12688/f1000research.2-152.v1)

Introduction

Gastrointestinal stromal tumors (GIST) are sarcomas arising in the muscle wall of the gastrointestinal tract. The majority of GISTs are driven by activating mutations in the receptor tyrosine kinases KIT or PDGFR- α , whose aberrant signaling induces uncontrolled proliferation and decreased apoptosis¹. Because of the small number of driving genetic mutations, GISTs represent a paradigm for kinase-driven solid tumors, and offer one of the best models to shed light on fundamental questions about cancer, providing a critical understanding of more genomically complex solid tumors.

Tyrosine kinase inhibitors (TKIs), such as imatinib and sunitinib, block the aberrant activation of KIT, leading to major clinical benefits of objective response and durable disease control. Imatinib represents the standard first-line therapy for this disease when it is surgically incurable. However, imatinib does not cure advanced GIST². Moreover, while the large majority of patients treated with imatinib do not show signs of primary resistance (disease progression within the first six months of therapy), secondary resistance to imatinib emerges in at least half the patients after two years of therapy and in more than 80% of patients after seven years^{1,3,4}.

Imatinib was originally introduced to treat chronic myeloid leukemia (CML), because of its specificity for the TK domain in the *bcr-abl* translocation gene, which characterizes the disease. Imatinib represents the standard first-line therapy and has led to dramatic improvements in outcomes for CML patients in the chronic phase: at six years the estimated event-free survival is 83%, and an estimated 93% of treated patients are free from progression⁵. Importantly, secondary resistance is far less frequent in CML⁵.

In this article we consider two fundamental, clinically relevant, but still unanswered questions: Why do patients with advanced GIST often relapse and does secondary resistance originate before or during TKI treatment?

Materials and methods

In order to obtain estimates for the probability of at least one drug resistant GIST cell being present at the time the tumor reaches a given diameter, and before the introduction of the TKI, we used formula (5) in Tomasetti *et al.*⁶:

$$P_R = 1 - e^{-uM \left(\frac{1-a/2-b}{1-a-b} \right) \frac{l(1-a-b)}{d+lb} \ln \left(\frac{l(1-a-b)}{l(1-a-2b)-d} \right)}$$

This formula estimates the probability of having resistant mutants P_R in a tumor of size M , (number of cells), and is derived by counting the number of divisions required for the tumor to reach that size. It is assumed that, at each cell division, there is a small probability u that one of the daughter cells is hit by a mutation known to induce drug resistance. The parameter l and d are the birth and death rates for the cell population, while a and b are the probabilities for the cells' mode's of division. This mathematical result is based on standard assumptions and has been successfully used to predict the development of acquired resistance to targeted epidermal growth factor receptor

(EGFR) blockade in colorectal cancer⁷. In order to apply this formula we used parameter estimates available in the literature, as follows.

The somatic point mutation rate u can be estimated to be between 10^{-9} and 10^{-8} per base per cell division^{8,9}. There are many known point mutations causing resistance to imatinib¹⁰. To obtain a conservative bound, we assumed a value of 10 point mutations in our calculations, though the actual number could be higher¹¹. Overall, the probability of a point mutation causing drug resistance in GIST should then be at least 10^{-8} per cell division. It has been estimated that 10^{-9} cancer cells are present per cm^3 of tumoral mass¹². As our goal was to generate an unfavorable scenario to our hypothesis, and since stromal tissue and other types of cells may be present, we halved this amount. Long-term drug resistance requires secondary mutations to be present in cells that are long-lived or able to self-renew¹³. KIT is known to have anti-apoptotic activity, thus contributing to cell lifespan¹⁴. Alternatively, resistance could arise from mutations in rare cells possessing stem cell attributes. Interstitial cells of Cajal progenitors are a potential candidate¹⁵. Their frequency is estimated to be 6.2×10^{-3} of all cells¹⁶. To be conservative, we only considered the stem cell compartment and set all other parameters of the main formula (5) in Tomasetti *et al.*⁶ to zero.

Using these results and parameters values, we can derive the lower bound of Figure 1 for the probability of at least one drug resistant GIST cell being present at the time the tumor reaches a given diameter, and before the introduction of the TKI. Because we selected parameters to generate an unfavorable scenario to our hypothesis, we expected the actual curve to be above the curve of Figure 1. The calculation and the figure were obtained using the freely available R software (version 2.15.3)¹⁷.

Results

We combined experimental data with our mathematical formulas⁶ that recently have been successfully used to predict the development of acquired resistance to targeted EGFR blockade in colorectal cancer⁷. By so doing, we have obtained the relationship shown in Figure 1 between the tumor diameter at detection, and the probability that the tumor already harbors a resistant mutant at that time (see Materials and methods for the derivation of Figure 1). For example, for GISTs with diameters of 2 and 6 cm we estimate this probability to be equal to 0.12 and 0.97, respectively.

We now bring these elements to bear in interpreting recent clinical observations. The size of a GIST at presentation may vary between 1 and >40 cm in diameter¹. For example, in a clinical trial (NCT00237185) of 147 patients with unresectable or metastatic GIST expressing KIT and treated with imatinib, 75% of the patients had a GIST whose diameter was larger than 7 cm at treatment¹⁸. Based on Figure 1, we would expect the large majority of these patients to have resistant mutant cells already present by the time of GIST detection and therefore to develop resistance during therapy. In fact, only about 20% of the patients in this clinical trial were still progression-free after five and a half years from the start of imatinib treatment¹⁸.

The combination of experimental and clinical data with our mathematical estimates implies then that secondary resistance in patients

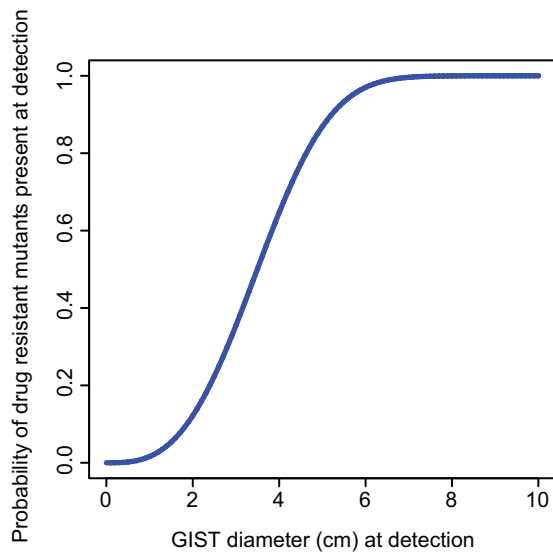


Figure 1. The estimated lower bound for the probability that mutant cells resistant to imatinib are already present in a gastrointestinal stromal tumor (GIST) at time of treatment, as a function of tumor diameter.

with advanced GIST is to be expected and is due to the large size of the tumor at detection time. That is, mathematical estimates indicate that mutations responsible for secondary resistance are already present before the start of the treatment in patients with large tumor sizes, a key factor in explaining the high relapse of advanced GIST patients to TKI.

Discussion

It has been recently suggested that secondary resistance may be the result of the treatment itself rather than pretreatment mutations⁴. It is worth considering this issue further. Secondary mutations are not usually found at detection using conventional Sanger sequencing techniques⁴. However, these techniques are not sensitive for rare events. Because imatinib has not yet selected for resistant mutants, they may remain extremely rare, and thus are likely missed by the assays used for mutation analysis. Thus, the fact that mutants are rarely found at detection does not contradict the possibility that the point mutations did occur before treatment. Using our mathematical formulas again⁴, we estimate that if resistance is present in a GIST of 2 cm in diameter, only approximately 1 out of 10^8 tumor cells will be drug resistant. Most importantly, while resistance may also originate during treatment, this does not exclude the fact that random point mutations can occur during the re-treatment phase at each cell division. We have shown that the estimated number of resistant mutants produced before the start of the treatment is sufficient to explain the observed clinical data. This logic also holds for the development of tertiary resistant clones that emerge within, on average, six months of starting second-line therapy with sunitinib following failure of imatinib¹⁹. Thus, our calculations, while not addressing the problem of primary resistance, are able to explain secondary resistance in advanced GIST.

Our prediction that patients with smaller GISTs at detection will have a smaller probability of progression has been already supported

by the clinical correlation of tumor size at presentation with improved outcomes on imatinib¹⁸. With sufficient data, it may also be possible to observe a relation between tumor diameter at detection and probability of resistance that is similar in shape to Figure 1, since a resistant clone originating before therapy should eventually grow to a detectable size. Lastly, in the future, with next-generation DNA sequencing techniques, it may become possible to detect the presence of secondary mutations that represent a very small fraction of cells at initial presentation.

Our results, if confirmed, would have important clinical implications. A therapy using a combination of TKIs (for example by adding a new agent with novel spectrum activity to imatinib) would select only for those cells which have been hit by two point mutations, each causing resistance to one of the drugs. We estimate that such a probability is very small for realistic values of the GIST diameter (not considering cross resistance). Therefore, combination therapy could drastically reduce the development of drug resistance caused by point mutations. Also, earlier detection of GIST offers the potential to provide measurable improvements in outcome, as long as it can result in a sufficiently large reduction in the average diameter at detection.

Author contributions

CT and GP conceived the idea. CT, GDD, and GP designed and performed research, and wrote the manuscript. CT contributed the mathematical analysis.

Competing interests

CT and GP have no competing interests to disclose. GDD discloses the following: Novartis. Consultant, <\$10k per annum. Honorarium <\$10k per annum. Research support to Dana Farber for specific clinical trial agreements in our sarcoma unit, >\$200,000 per year. Research support to Dana-Farber for specific clinical trial agreements in our sarcoma unit, >\$200,000 per year. Pfizer. Consultant, <\$10k per annum. Honorarium <\$10k per annum. Research support to Dana-Farber for specific clinical trial agreements in our sarcoma unit, >\$200,000 per year. Infinity Pharmaceuticals. Consultant, <\$10k per annum. Research support to Dana-Farber for specific clinical trial agreements in our sarcoma unit, >\$200,000 per year. Regulatory presentation support, uncompensated. Glaxo Smith Kline. Consultant, <\$10k per annum. Kolltan Pharmaceuticals. Chair, Scientific Advisory Board. Chair, Medical Advisory Board. Consultant. Equity (minor stake, non-public).

Grant information

The work of CT was supported in part by the National Institute of Health under Grant T32 CA009337. GDD was partially funded by DFHCC SPORE (NCI Grant) for GIST research, as well as Ludwig Center for Cancer Research at Dana-Farber/Harvard Cancer Center. GP was supported in part by the NIH/NCI grant 5P30 CA006516-46.

The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

References

1. Corless CL, Heinrich MC: **Molecular pathobiology of gastrointestinal stromal sarcomas.** *Annu Rev Pathol.* 2008; **3**: 557–86.
[PubMed Abstract](#) | [Publisher Full Text](#)
2. Le Cesne A, Ray-Coquard I, Bui BN, *et al.*: **Discontinuation of imatinib in patients with advanced gastrointestinal stromal tumours after 3 years of treatment: an open-label multicentre randomised phase 3 trial.** *Lancet Oncol.* 2010; **11**(10): 942–9.
[PubMed Abstract](#) | [Publisher Full Text](#)
3. Antonescu CR, Besmer P, Guo T, *et al.*: **Acquired resistance to imatinib in gastrointestinal stromal tumor occurs through secondary gene mutation.** *Clin Cancer Res.* 2005; **11**(11): 4182–90.
[PubMed Abstract](#) | [Publisher Full Text](#)
4. Antonescu CR: **The GIST paradigm: lessons for other kinase-driven cancers.** *J Pathol.* 2011; **223**(2): 251–61.
[PubMed Abstract](#) | [Publisher Full Text](#)
5. Hochhaus A, O'Brien SG, Guilhot F, *et al.*: **Six-year follow-up of patients receiving imatinib for the first-line treatment of chronic myeloid leukemia.** *Leukemia.* 2009; **23**(6): 1054–61.
[PubMed Abstract](#) | [Publisher Full Text](#)
6. Tomasetti C, Levy D: **Role of symmetric and asymmetric division of stem cells in developing drug resistance.** *Proc Natl Acad Sci U S A.* 2010; **107**(39): 16766–71.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
7. Diaz LA Jr, Williams RT, Wu J, *et al.*: **The molecular evolution of acquired resistance to targeted EGFR blockade in colorectal cancers.** *Nature.* 2012; **486**(7404): 537–40.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
8. Araten DJ, Golde DW, Zhang RH, *et al.*: **A quantitative measurement of the human somatic mutation rate.** *Cancer Res.* 2005; **65**(18): 8111–7.
[PubMed Abstract](#) | [Publisher Full Text](#)
9. Drake JW, Charlesworth B, Charlesworth D, *et al.*: **Rates of spontaneous mutation.** *Genetics.* 1998; **148**(4): 1667–86.
[PubMed Abstract](#) | [Free Full Text](#)
10. Heinrich MC, Corless CL, Blanke CD, *et al.*: **Molecular correlates of imatinib resistance in gastrointestinal stromal tumors.** *J Clin Oncol.* 2006; **24**(29): 4764–74.
[PubMed Abstract](#) | [Publisher Full Text](#)
11. Wang WL, Conley A, Reynoso D, *et al.*: **Mechanisms of resistance to imatinib and sunitinib in gastrointestinal stromal tumor.** *Cancer Chemother Pharmacol.* 2011; **67**(Suppl 1): S15–24.
[PubMed Abstract](#) | [Publisher Full Text](#)
12. Klein M, Bartoszynski R: **Estimation of growth and metastatic rates of primary breast cancer.** In *Mathematical Population Dynamics.* (eds. Arino, O., Axelrod, D.E. & Kimmel, M.) (Marcel Dekker, New York, 1991).
13. Tomasetti C: **A new hypothesis: imatinib affects leukemic stem cells in the same way it affects all other leukemic cells.** *Blood Cancer J.* 2011; **1**(5): e19.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
14. Rossi F, Ehlers I, Agosti V, *et al.*: **Oncogenic Kit signaling and therapeutic intervention in a mouse model of gastrointestinal stromal tumor.** *Proc Natl Acad Sci U S A.* 2006; **103**(34): 12843–8.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
15. Bardsley MR, Horvath VJ, Asuzu DT, *et al.*: **Kitlow stem cells cause resistance to Kit/platelet-derived growth factor alpha inhibitors in murine gastrointestinal stromal tumors.** *Gastroenterology.* 2010; **139**(3): 942–52.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
16. Lorincz A, Redelman D, Horvath VJ, *et al.*: **Progenitors of interstitial cells of cajal in the postnatal murine stomach.** *Gastroenterology.* 2008; **134**(4): 1083–93.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
17. **R_Core_Team. R: A Language and Environment for Statistical Computing.** 2.15.3 edn (R Foundation for Statistical Computing, Vienna, Austria, 2013).
[Reference Source](#)
18. Blanke CD, Demetri GD, von Mehren M, *et al.*: **Long-term results from a randomized phase II trial of standard- versus higher-dose imatinib mesylate for patients with unresectable or metastatic gastrointestinal stromal tumors expressing KIT.** *J Clin Oncol.* 2008; **26**(4): 620–5.
[PubMed Abstract](#) | [Publisher Full Text](#)
19. Demetri GD, van Oosterom AT, Garrett CR, *et al.*: **Efficacy and safety of sunitinib in patients with advanced gastrointestinal stromal tumour after failure of imatinib: a randomised controlled trial.** *Lancet.* 2006; **368**(9544): 1329–38.
[PubMed Abstract](#) | [Publisher Full Text](#)

Current Referee Status:

Referee Responses for Version 1



Gerald Prager

Department of Medicine, University of Vienna, Vienna, Austria

Approved: 18 July 2013

Referee Report: 18 July 2013

Tomasetti *et al.* were combining clinical and experimental observations with mathematical formulas to estimate that in advanced GIST, the genetic changes responsible for resistance are already present from the beginning of treatment. The model is of highest interest, especially in respect of second and third line treatment with targeted therapies. However, prospective validation is required before this model can be incorporated in combination drug trials.

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Competing Interests: No competing interests were disclosed.



Shiro Urayama

Department of Internal Medicine, University of California Davis Medical Center, CA, USA

Approved: 18 July 2013

Referee Report: 18 July 2013

This is a brief report depicting a mathematical model based estimation of presence of chemoresistant cells in gastrointestinal stromal tumor (GIST). The model incorporates the notion of the presence of rare cells possessing stem cell attributes (cancer stem cells) among heterogeneous cells within such mass, contributing to the chemotherapeutic resistance. On the basis of genetic point mutation as a source for the resistance, the probability formula for existence of resistant cells at a tumor size was constructed. Despite some constraints, the article presents reasonable assumptions providing for the lower bound of the probability. Based on this model, many of the clinical index GIST mass has significant likelihood of containing a tyrosine kinase inhibitor resistant cell at the time of the discovery. If future clinical findings support the assumptions made, the model would demonstrate a rationale for combination therapeutics.

Specific points:

Title and Abstract: Appropriate

Article content: Methods and approaches are described appropriately and references made to the main formula utilized

Conclusions: Appropriate

Data: Appropriate as included

Minor item:

Materials and Methods second paragraph, 5th sentence should have “10⁹ cancer cells” instead of “10⁻⁹ cancer cells are present per cm³ of tumor mass”.

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Competing Interests: No competing interests were disclosed.

1 Comment

Author Response

Cristian Tomasetti, Johns Hopkins University, USA

Posted: 11 Nov 2013

We thank the referee. It should read "*estimated that 10⁹ cells*".

Competing Interests: I am Cristian Tomasetti, one of the authors of the article.