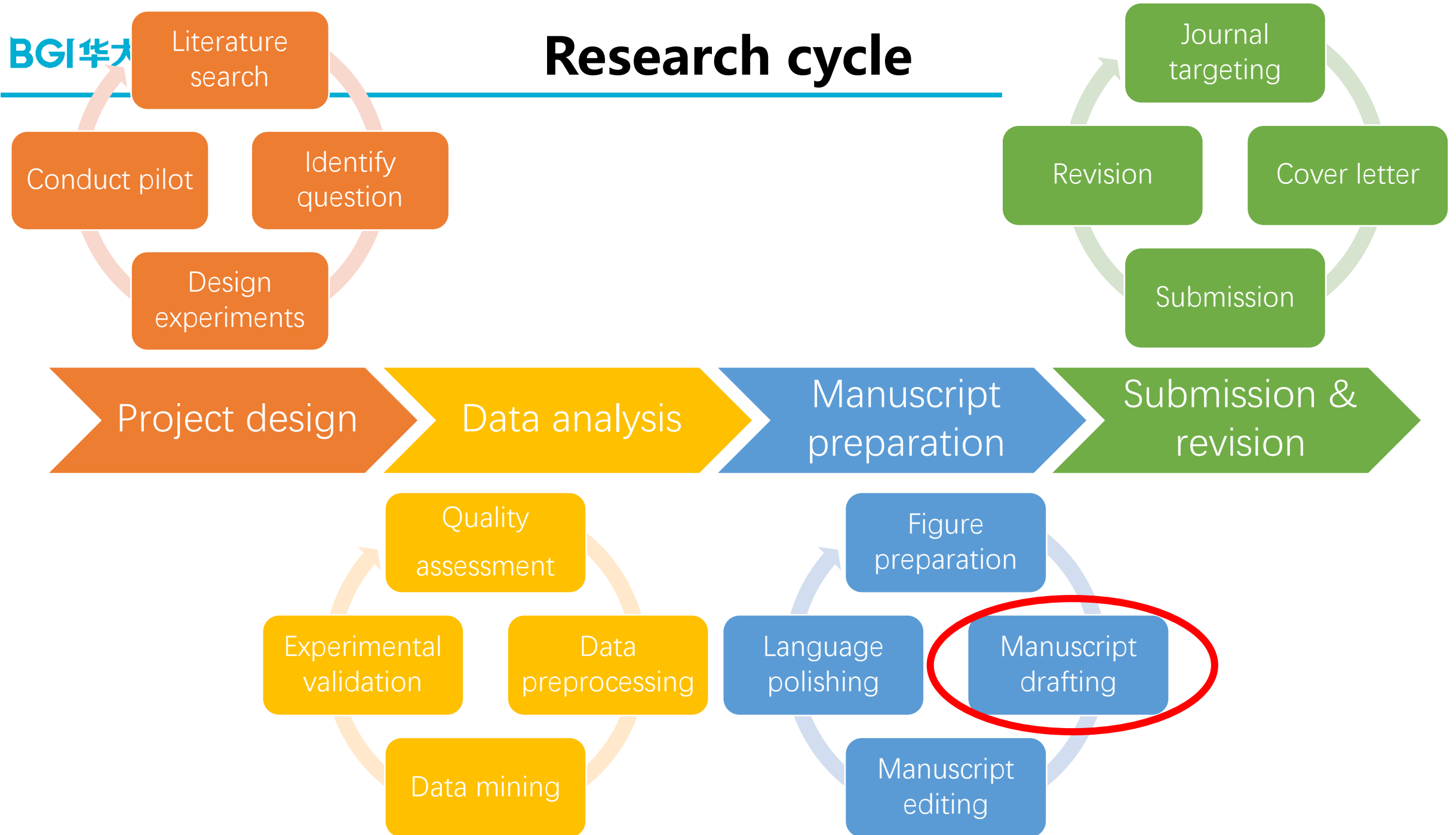


Scientific Writing

魏桐

4/26/2023

Research cycle



What is scientific writing

- Scientific writing is **a technical form** of writing that **communicates** scientific concepts/findings to other scientists in a written form.
- Depending on the specific scientific genre—a journal article, a scientific poster, or a research proposal, **the details vary but have a greater similarity**.
- Important hallmarks of all scientific writing are,
 - Has to be **logical and organized**,
 - Must be **exact and precise**,
 - Needs to be set **within the context** of published work.

How to get your work published

- Your research is good...
 - It has novelty.
 - The experiments is well-designed.
 - The results are solid.
 - ...
- And more importantly nowadays, your manuscript is **well organized and written!**

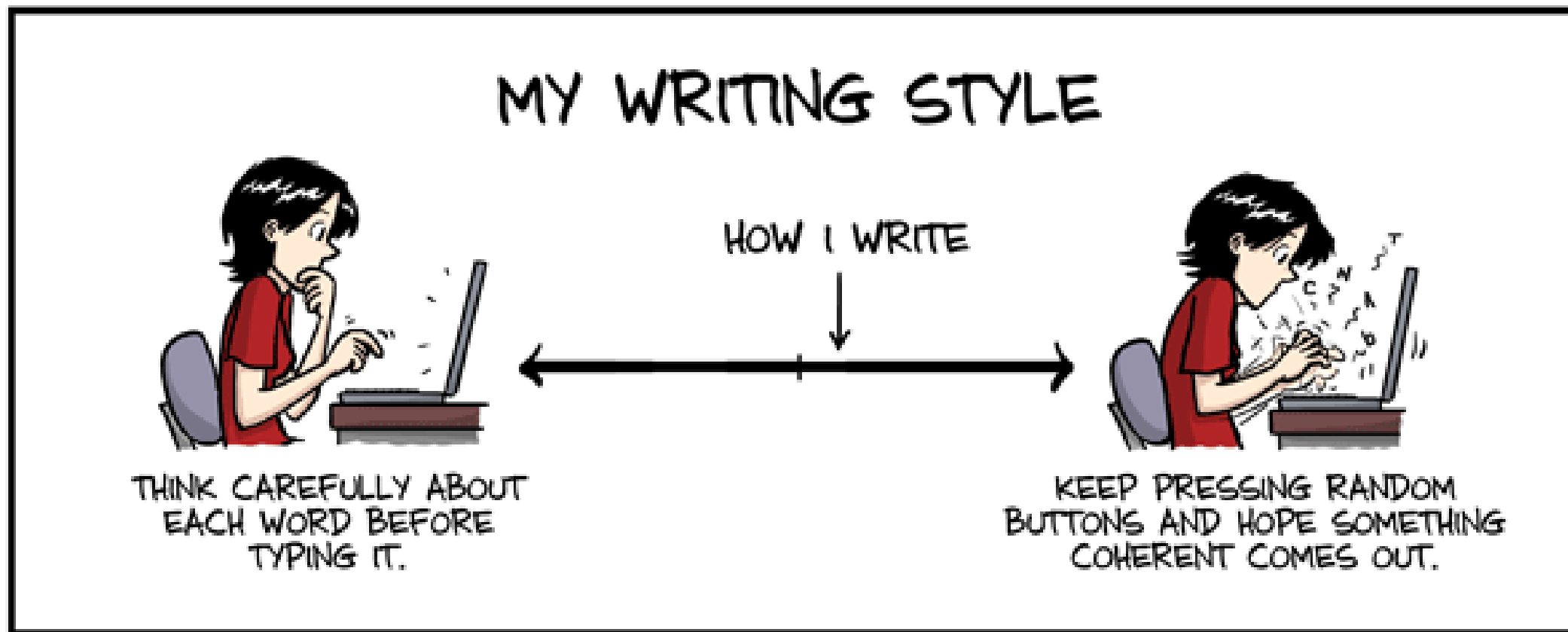
What happens to poorly written papers

- Editors **miss** the point.
- Editors feel the paper **will not pass** the reviewing process.
 - Most likely reject.
- Reviewers feel **frustrated**.
 - Likely reject.
- Reviewers feel **confused**.
 - Probably raise extra questions or ask for more results.

Writing a manuscript is
as important as doing experiments.

For high-impact journals,
writing is often more important.

Writing needs practices



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Key questions related to your research

- **Why** did you do it?
- **How** did you do it?
- **What** did you get?
- **So what** did it mean?

Q&As related to your work

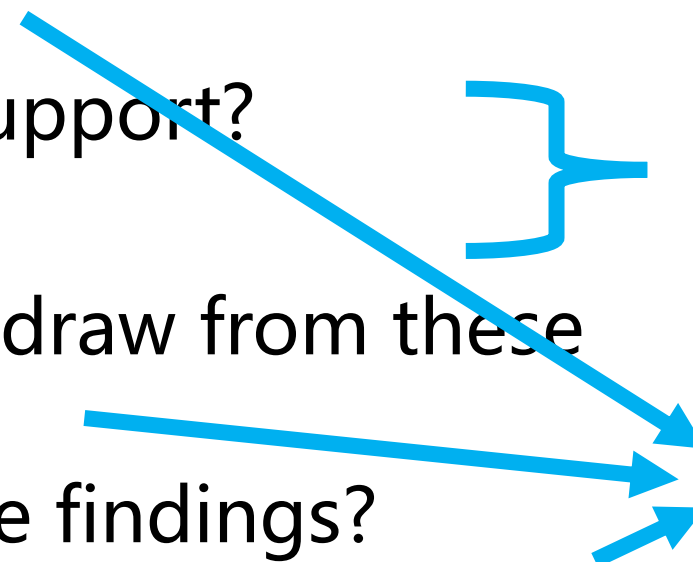
- Why is your work important?
- What problem do you want to solve?
- What is your main discovery?
- What experiments did you do to support?
- How did you do the experiments?
- What specific conclusions can you draw from these experiments?
- What can you conclude from all the findings?
- What is the significance of your discovery?

} WHY

} HOW

WHAT

SO WHAT



Abstract

- Background + question
- A sentence or phrases about methods
- Major discoveries
- Conclusion + significance

WHY



Introduction

HOW



Methods

WHAT



Results

SO WHAT



Discussion

- Why did you do it?
 - ... is important in organ development/crop science; however, the mechanism remains unclear.
- How did you do it?
 - We carried out a multi-omic approach ...
- What did you get?
 - The results showed that genes were associated with ...
- So what did it mean?
 - Our work discovered key players and shed light on ...

Part I

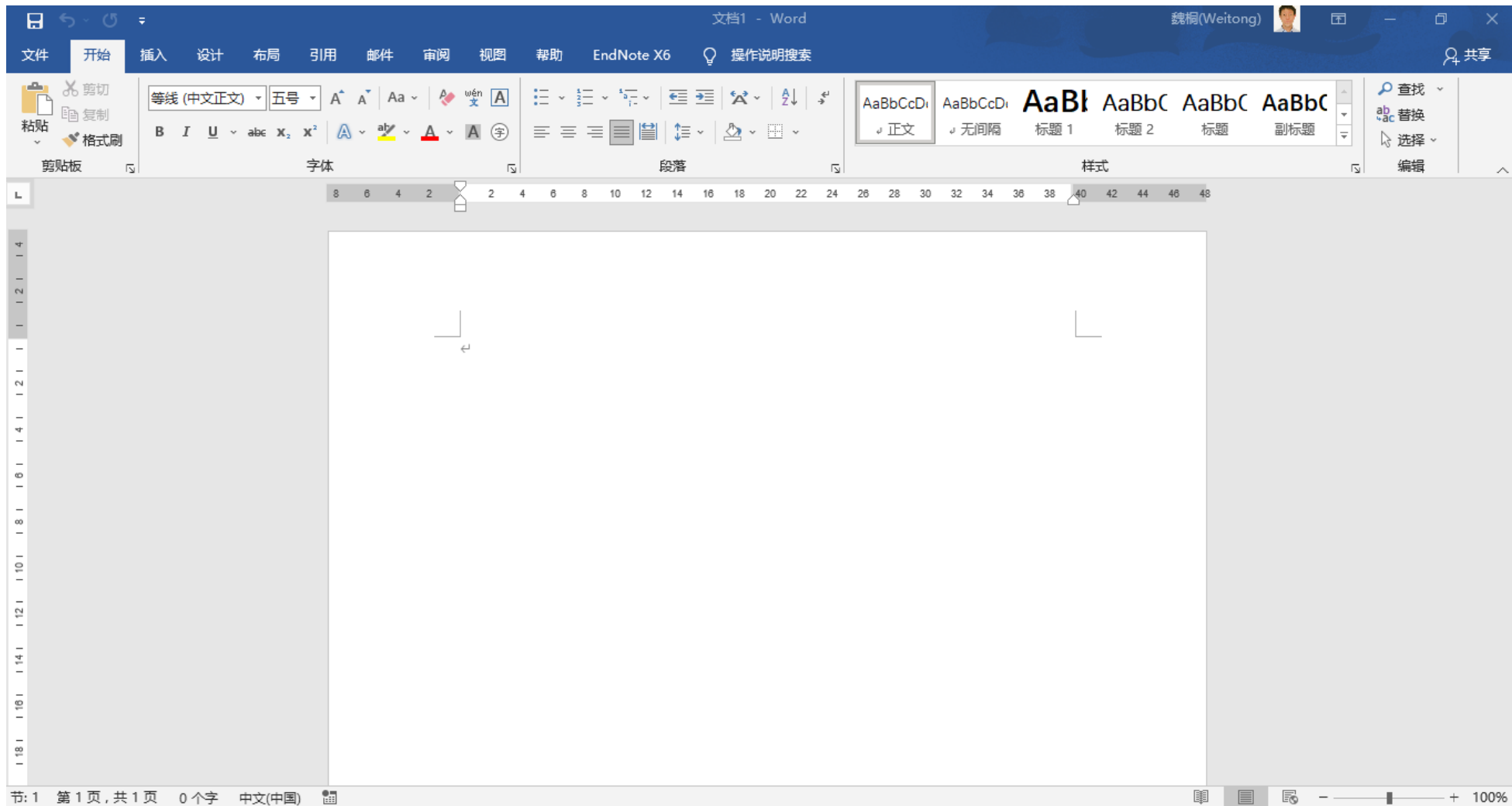
Manuscript drafting

3 phases during manuscript preparation

- The preparation phase
 - A **final set of processed data**, assembly, variants, expression, etc.
 - A complete set of **preliminary figures & tables**
- The drafting phase
 - **An story line** with logic
 - Figures + Results & Methods
 - Discussion + Introduction
 - Abstract + Title
- The editing phase
 - Seek for **professional comments**
 - Language polishing

- Get all the results together, e.g. processed results (**figures & tables if possible**)
 - Get the methods ready, i.e. software, parameters, filtering criteria, etc.
 - Scan the literature briefly
 - Revisit **the scientific question(s)**
-
- Conceptualize/visualize your work, and draw the OUTLINE

Ready to draft the manuscript!



- **OUTLINE** guides the drafting
- **BOTTOM-UP** writing
- Write out **an rough Abstract** from the outline
- Assign **a topic sentence** for each paragraph in each section
- Image **writing for people** without any background knowledge

Outcomes from the drafting phase

- **An rough abstract** with logic
- Main figure layout
- **Results & Methods**
- A rough Discussion
- A rough Introduction
- A revised Abstract if results change
- A working title

- It answers the **WHY, HOW, WHAT** and/or **SO WHAT** questions.
- It is a special tool to,
 - Guide figure preparation,
 - Dictate results writing,
 - Review the whole draft.
- **Know your data** and spent some time, say 30 minutes, to draw the outline.

- Describe all the experimental procedures **in details**
 - Experimental materials, species, ecotypes, batches, etc.
 - Treatment, chemicals, concentration, time, etc.
 - Data analysis, software versions, parameters, formats, etc.
- Write methods **after the analyses is done**.
- This is quite straightforward.

- Layout figures in **a logical order**
 - Combine the figures **related to the same topic**
 - Write the simple legends
 - Expand into **structured** results
-
- Write **section by section** if possible
 - Write panel by panel if facing some difficulty

- Make them **self-explanatory**
 - People do not need to read legends to understand
 - Be informative and be pretty
 - State findings in figure titles
 - Explain everything in legends
-
- Learn from good journals
 - Avoid use red & green

Tips for writing result sections

- Each paragraph should have **ONLY one topic sentence**.
- Paragraphs should be **logically organized** around headings or sub-headings.
- Do not count words closely.

Discussion example: questions answered

WHY

WHAT

As germplasms of major crops are maintained as genebank collections, understanding the population structure and phylogenetic relationships is of great importance for genebank management and utilization. In lettuce breeding, the GP1 species are used widely as there is no reproductive barrier within the group^{6,7}. Our phylogenetic analyses clarified several issues regarding the taxonomic status of these GP1 species (see the Supplementary Note for a detailed discussion). First, the presumed GP1 species *L. georgica* should be reassigned as it clustered with the GP3 species *L. virosa*. The *L. dregeana* and *L. sagittata* samples are not to be considered as true wild species. Another GP1 species, *L. altaica*, has been considered as conspecific with *L. serriola*^{29,30}, but the plastid phylogeny implied an introgression and fixation of a distantly related plastid haplotype in *L. altaica*. Phylogenetic analyses with additional samples will clarify these taxonomic issues in wild species. Our study also pointed out future directions in germplasm collection and utilization. Among the investigated samples, *L. serriola* from the Caucasus represents the most promising resource because the population from this area showed the highest nucleotide diversity. *L. aculeata* represents another potentially important gene pool, as its phylogenetic position distinct from other GP1 species suggests a different genetic repertoire. Thus, our study provided new insights regarding accession identity and genetic resources for crop improvement, demonstrating the value of whole-genome sequencing in the management of crop collections and the utilization thereof.

HOW

SO WHAT

Personal note

- Background information
 - Describe **the importance** of your area of study
 - Review the major findings in the field
 - Introduce **the related concepts/pathways/genes**
- Scientific question(s)
 - Raise the question **in a logical way**
 - Stress **its significance**
- Your findings
 - Communicate the major discovery

- The space is limited, normally 1-3 paragraphs.
 - Use only materials present **in the literature**
 - Use only materials **directly related** to your work
- Keep writing
 - Use your background knowledge
 - Do not overthink on details
 - Add (ref/cite) for important information and prepare a reference file

- It should emphasize information on the **MAIN concept/discovery** you want to tell readers.
 - What was **DONE**
 - What was **FOUND**
- It is not easy to find a precise title, but you should have after the manuscript is drafted.

- **TOP-DOWN** editing
- Read through the draft multiple times
- Do not try to fix everything once; fix on **one thing/error at a time** (tense, details, typo, etc.)

The editing phase (continued)

- Edit structure **section by section**
 - Move sentences/paragraphs to where they belong
 - Focus on the logic in each section
- Edit paragraph **line by line**
 - Look into each sentence, word, phrase, etc.
 - Language skills play an important role
 - Do it several rounds
- Seek for **professional feedbacks** after you have done a thorough editing
- Final check in spelling, grammar, numbers, figures & tables, etc.

Outcomes from the editing phase

- **Clear key questions**
- The main Figures and Results **in the same topic**
- An updated Discussion and Introduction **after a thorough review of literature**
- A rewritten Abstract with **answers to the 4 questions**
- A precise and interesting Title
- Methods, supplementary materials, data deposition, etc.

- That (referring to a specific thing) vs which (adding a clause)
- Correct/incorrect, not right/wrong
- Only use “significant” for statistics
- Give exact/approximate numbers instead of several, most, a few, etc.

Tips for English writing (continued)

- Articles: a, an or the
- Tense: past tense for results; present for facts; present tense in figure legends and formula; present perfect tense for continuous work
- Could, may, might
- Demonstrate, indicate, suggest
- Use short sentences

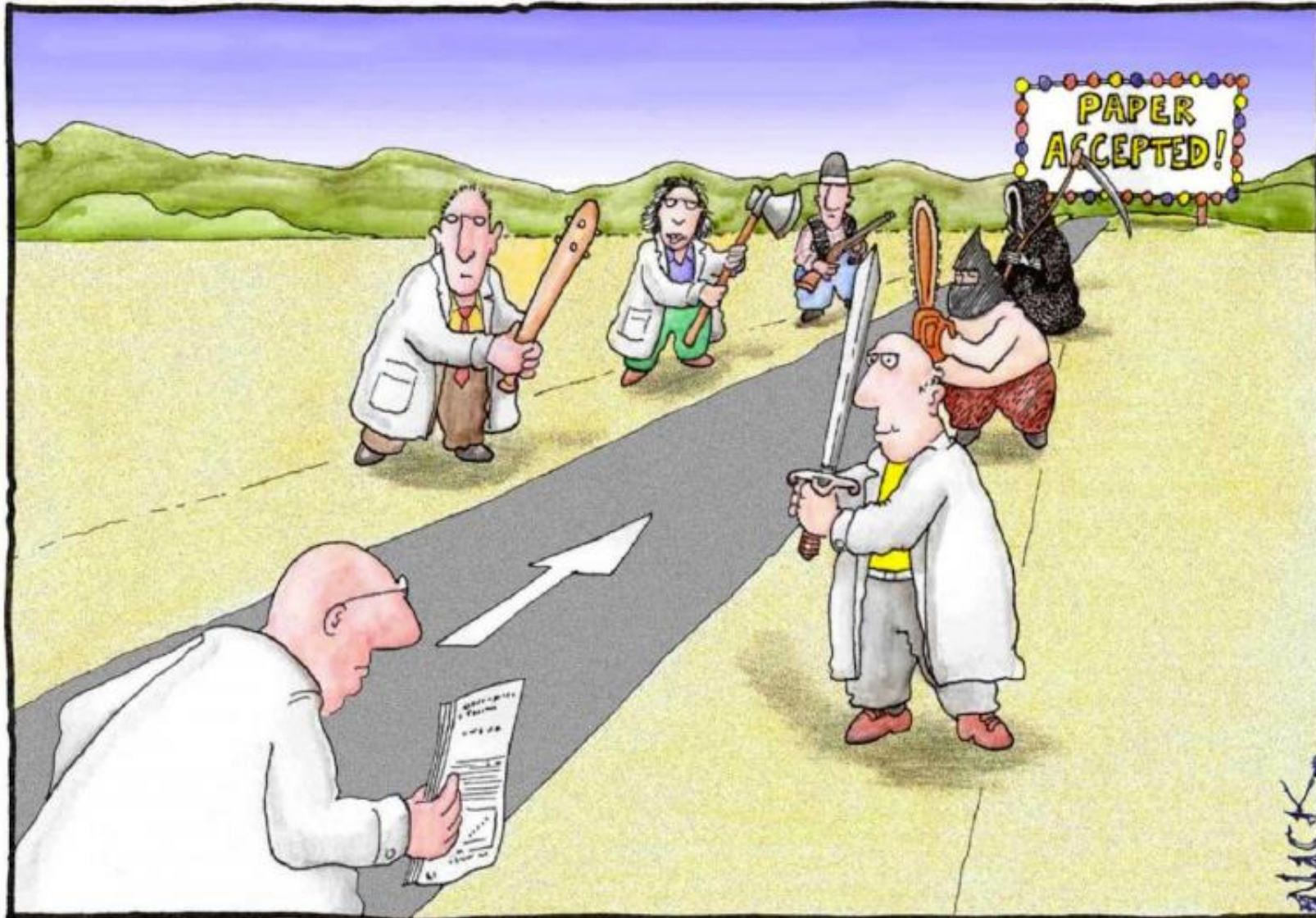
- Check grammar!

- Abstract is the essence of your manuscript
 - It is **the FIRST thing** readers see, and sometimes the ONLY thing
 - It **sets the stage** by telling what your work is about
 - It is **critical for the reviewing process**
- It should be one paragraph about,
 - Why you did the work
 - What you found
 - Why it matters

Part II

Submission & revision

Ready to submit your manuscript!



There is **no end of editing**, but there should be **a deadline for submission!**

- Everyone is satisfied (seldom).
- Major authors are satisfied (sometimes).
- Major authors are tired (most of the time).
- My advice
 - All the **major issues** raised by professionals are solved.
 - The logic is **easy to follow**.
 - The main text **reads well**.
 - There will not be any significant improvement in the next 2 months.

- Before pushing the SUBMIT button
 - Choose a **reasonable** journal
 - Write a **thoughtful Cover Letter**
 - Reformat according to the guideline
- Major or minor (unlikely) revisions
- Conditional acceptance -> copyediting -> formal acceptance

- Impact Factor
 - It shows how many times **on average** papers in one year have been cited **over the next two years**
 - The top journal is not always a good choice
- The potential readers
 - Make a list of journals with narrow to broad readership
- Open access
 - More read and cited
 - Required by funding agencies

Choose a target journal (continued)

- Time to publication
 - Competition
 - Graduation
 - Annual report
 - etc.

Things to do for a particular journal

- **A thoughtful cover letter**
 - to communicate with editor
- A well-written abstract
 - Answer the WHY, HOW, WHAT, and SO WHAT questions
- The checklist
 - Follow the author guideline

- Cover letter is for editors, who decide whether to send out your manuscript for reviewing.
- Do **NOT** copy Abstract. **ALWAYS** write a good letter!
 - One page
 - Start with a small paragraph about the paper you are submitting
 - Explain why it is **important and interesting**
 - Describe what you have concluded
 - Describe how it **fits into the scope**

With our best regards,
Corresponding authors

Compared to their economic and biological importance, viruses are very much under-investigated. Economic and genetic resources are needed. Genetic improvement of crops for better resistance to virus is also inseparable with other species in the

caranther genome by a novel combination of the de novo-generation Illumina GS sequencing technology, the 350Mb genome and ~99% of the gene regions. The abundance of satellite sequences and HSA4. Caranther (8,022) in Arabidopsis, but no other whole genome wide random gene duplications were observed among the *Slac* genome sequenced. Nevertheless, tandem duplications genes related to caranther biology, such as myosin cytoplasmic II, secondary antibodies that give caranther and bitter taste, and oil seed modification associated with the myosin and vasculature genes map to the caranther of the 7 caranther chromosomes were result of false of association with pairs. As predicted, most caranther and by ordered on the chromosome with 99% sequence

cucumber genome was affordable it is only when semi-genome sequencing is come to stage. So genomics is the 'equation', eliminating or narrowing the two model organisms and orphan species. This research signals the arrival of any changes to genetic improvement of orphan crops, which is still a critical sustainable development of the rural disadvantaged regions in the world.

For more information, contact the author.

- Format the manuscript according to the guideline
 - Word count (make each section close to the requirements)
 - Text formatting (title page, subtitles, spacing)
 - Figure formatting (citation, width, etc)
 - Supplementary materials

PRINT PAPER

LETTER

Outgrowth of single oncogene-expressing cells from suppressive epithelial environments

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[illegible]

The outgrowth of synaptic axons onto cortical target cells is regulated by a fundamental in vitro mechanism. However, synaptic direction and efficiency are also regulated by extracellular cues that are cell type- and synapse-specific (1,2), particularly with respect to the timing of synapse formation. Laminar and molecular gradients in the developing cortex regulate the timing of synapse formation (3,4). The synaptic outgrowth of retinal ganglion cell (RGC) axons onto cortical target cells is regulated by a number of extracellular cues, including laminar position (5), cell type (6), and synapse type (7). RGC axons in a transgenic mouse lacking retinotectal ephrins exhibit a dramatic reduction in the number of synapses formed, a failure to form synapses on nonacetylated nicotinic receptors (8), and a loss of their characteristic synaptic orientation (9). The synaptic outgrowth of retinal ganglion cell axons is affected by many additional extracellular cues

In order to acquire a high ΔG growth rate in the *in vitro* system, we first optimized the reaction temperature. At 37°C, the initial velocity of the reaction was 0.035 units/min, which was 10% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.05 units/min at 40°C, which was 14% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.06 units/min at 45°C, which was 17% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.07 units/min at 50°C, which was 20% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.08 units/min at 55°C, which was 23% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.09 units/min at 60°C, which was 26% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.10 units/min at 65°C, which was 29% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.11 units/min at 70°C, which was 31% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.12 units/min at 75°C, which was 34% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.13 units/min at 80°C, which was 37% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.14 units/min at 85°C, which was 40% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.15 units/min at 90°C, which was 43% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.16 units/min at 95°C, which was 46% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.17 units/min at 100°C, which was 49% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.18 units/min at 105°C, which was 51% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.19 units/min at 110°C, which was 54% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.20 units/min at 115°C, which was 57% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.21 units/min at 120°C, which was 60% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.22 units/min at 125°C, which was 63% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.23 units/min at 130°C, which was 66% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.24 units/min at 135°C, which was 69% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.25 units/min at 140°C, which was 71% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.26 units/min at 145°C, which was 74% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.27 units/min at 150°C, which was 77% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.28 units/min at 155°C, which was 80% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.29 units/min at 160°C, which was 83% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.30 units/min at 165°C, which was 86% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.31 units/min at 170°C, which was 89% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.32 units/min at 175°C, which was 91% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.33 units/min at 180°C, which was 94% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.34 units/min at 185°C, which was 97% of the maximal velocity (0.35 units/min) (Fig. 1). The initial velocity of the reaction was 0.35 units/min at 190°C, which was 100% of the maximal velocity (0.35 units/min) (Fig. 1).

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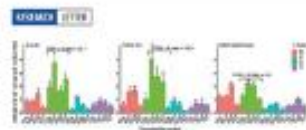


Figure 1. Silencing of human Borna disease virus (BDV) expression in HEK293 cells. Analysis of the effect of A. bisulphuratus on BDV and on the expression of the BDV genome (BDVg) in HEK293 cells. HEK293 cells were transfected with BDVg and A. bisulphuratus was added to the culture medium. The cells were harvested at 24, 48, and 72 h postinfection. The amount of BDVg in the culture medium was determined by quantitative real-time PCR. The results are expressed as the mean \pm SD of three independent experiments. * $P < 0.05$ compared with the control. The data are shown in Table 1.

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On the Cover: Authors, along with an additional Editorial Oversight team, submitted the *in situ* studies of the environment of the highly reduced pyrite in the Dabie orogen, as part of the special issue.

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[illegible]

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Key features:

- Includes main text and figures/tables/boxes plus references.
- Appears in print and online as a PDF and as full-text HTML.
- Text and figures are copy-edited to *Nature* style.
- Figure formats: colour artwork supplied in **RGB (recommended) or CMYK formats**; preferred formats are layered Photoshop (PSD) or TIFF for photographic images (minimum 300 p.p.i.), AI, Postscript, Vector EPS or PDF for line drawings and graphs.

Formatting details in text

Order of elements

Articles should be **ordered in the sequence:** title, authors, affiliations (plus present addresses), bold first paragraph, main text, references, tables, figure legends, (online-only) Methods (plus any associated references; data and code availability statements included at end of online Methods), acknowledgements, author contributions, competing interest declaration, additional information (containing supplementary information line (if any) and corresponding author line), Extended Data figure legends and Extended Data table titles and footnotes (any references unique to the Extended Data should be added to the end of the online-only reference list).

Fonts

We prefer the use throughout of a 'standard' font, **preferably 12-point Times New Roman**. For superscripts or subscripts, please apply actual super/subscript format; do not use 'raised' or 'lowered' formats. For mathematical symbols, Greek letters, and other special characters, use 'insert', 'symbol' and then select '(normal text)' or 'symbol' as the font. Use of other fonts can cause translation problems. List non-standard keyboard symbols in the letter accompanying the final accepted version of your paper.

Final print-only artwork

When preparing figures, authors are advised to refer to printed copies of Nature to get a sense of general size and style points. For an illustrated guide to preparing production-quality artwork after acceptance, see [this information document](#).

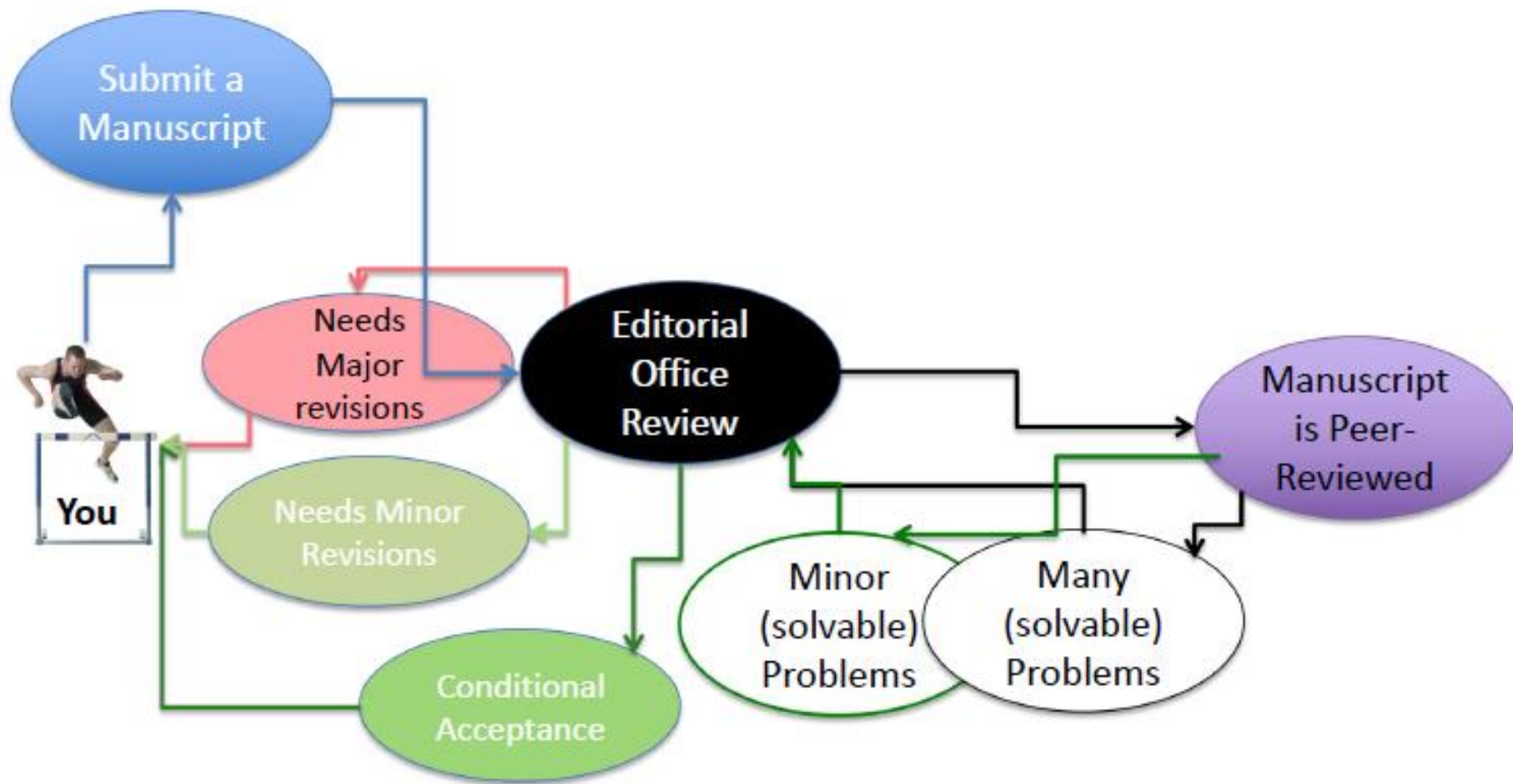
Lettering

Lettering should be in a sans-serif typeface, preferably Helvetica or Arial, the same font throughout all figures in the paper. Units, capitalization, etc. should follow Nature style. Where practical, avoid placing lettering directly over images or shaded areas.

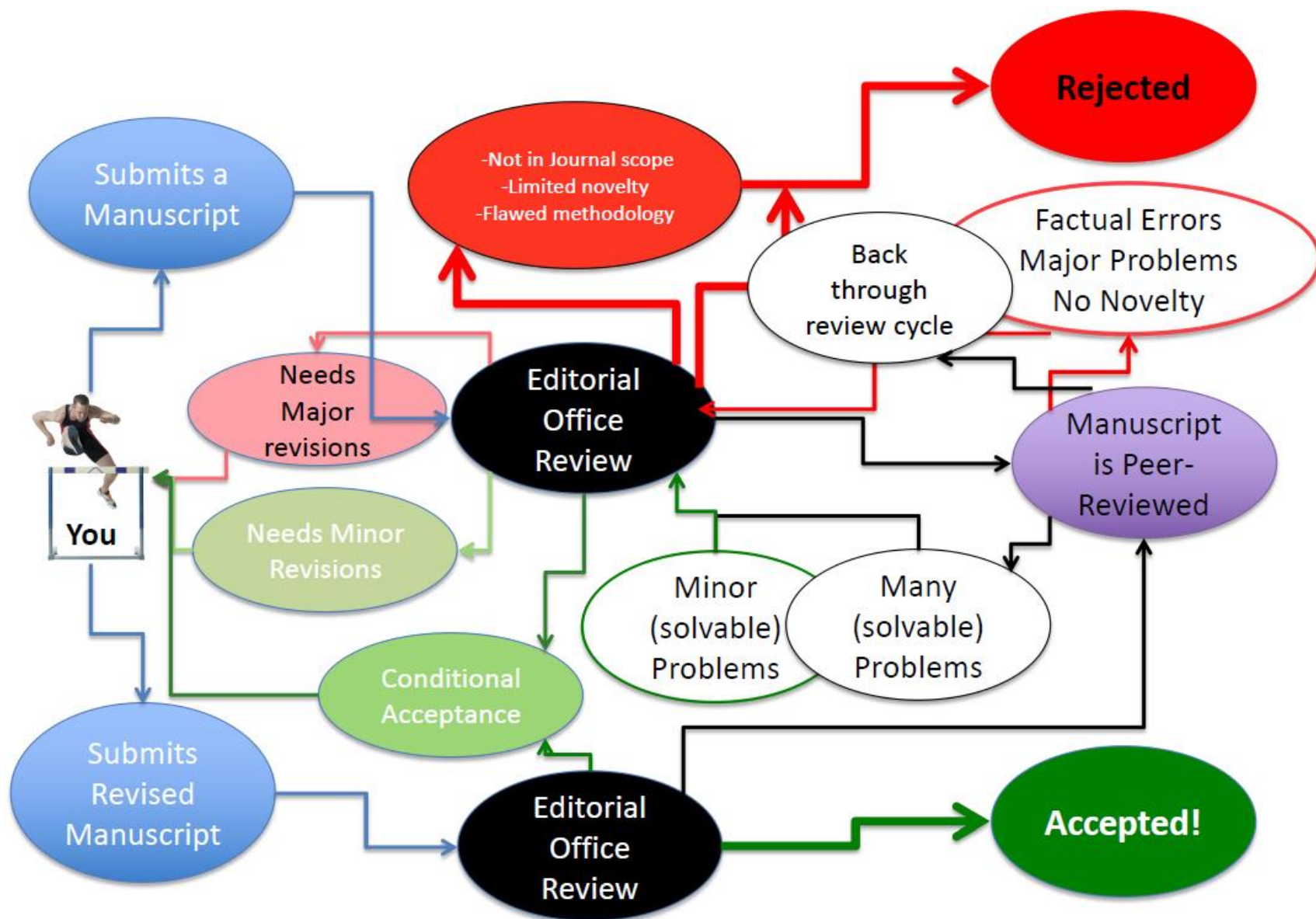
Separate panels in multi-part figures should each be labelled with **8 pt bold, upright** (not italic) a, b, c. **Maximum text size for all other text should be 7 pt; minimum text size should be 5 pt.** Amino-acid sequences should be presented in one-letter code in Courier.

Do not rasterize or covert text to outlines.

Theoretical reviewing process



Actual reviewing process



Common reasons for rejections

- No novelty
- Data do not support conclusions
- Not significant enough for your target journal
- Inappropriate experimental setup
- Inappropriate methods for the analysis
- Plagiarism
- Lack of ethical approval or missing data

Negative comments from reviewers

- Novelty
 - “There is very little biological novelty”
 - “The work is unlikely to have broad interest”
- Scientific questions
 - “Paper is mostly descriptive”
- English writing
 - “The English is difficult to follow”
 - “The methods are uninterpretable”

- Make a plan on,
 - What experiments/analyses to be done
 - What mistakes/typos need to be corrected
 - Which parts need to be re-written
 - Which comments need to be argued
- Focus on **the major issues**
- Fix the minor ones
- Highlight the changes in MS and indicate the line numbers in Response

Tips for response to reviewers

- Thumb of rules
 - **Respond politely**
 - Do NOT argue; take it easy
 - Response to EVERY issue
 - Provide new evidence if you did what the reviewers asked
 - Explain clearly the reason if you decide not to do
- It is difficult to argue with No Novelty
 - Highlight the novelty in the first submission
 - Point out the novelty if reviewers miss it

- Fill out required forms, author consent letter, checklist, statements, etc.
- Meet the requirements for final submission, word count, figure size, acknowledgements, etc.
- **Data and code availability**
- There is another round of copyediting.

Questions?