MRS is the world’s largest association for people and organisations that provide or use market, social and opinion research, business intelligence and customer insight.
Table of Contents

Introduction 4

The principles of the MRS Code of Conduct 5

Definitions 6

1: The Interviewer and the Participant 9

2: Meeting the Objectives 14

3: Structuring the Questionnaire 16

4: Writing the Questionnaire 18

5: Questionnaire Instructions 21

6: Piloting 22

7: Dealing with the Results 23

8: Further references 24

ANNEX: Ten Tips for DIY Surveys 25
Introduction

These Guidelines interpret the MRS Code of Conduct (revised 2014) and provide additional best practice guidance. Unless otherwise stated, Guidelines are not binding.

Research is founded upon the willing co-operation of the public and of business organisations. It relies on the confidence of those involved that it is conducted honestly, objectively, without unwelcome intrusion and without harm to participants. Its purpose is to collect and analyse information and not to create sales or to influence the opinions of anyone participating.

The general public and other interested parties are be entitled to complete assurance that every research project is carried out in accordance with the MRS Code of Conduct and that their rights and privacy are respected.

Rules from the MRS Code of Conduct applicable in each section of this document are stated in bold. These rules are binding on MRS members and MRS Company Partners and breaches may result in disciplinary action. The guidance that follows the rules provides interpretation and additional best practice. Members and Company Partners are reminded that this document is designed to complement the MRS Code of Conduct and should not be consulted in isolation.

As specified in the MRS Code, it is the responsibility of the researcher to keep abreast of any legislation which could affect research with employees and to ensure that all those involved in a project are aware of and agree to abide by the MRS Code of Conduct.

This material is provided for information only. It is not legal advice and should not be relied upon as such. Specific legal advice should be taken in relation to specific issues.
The Principles of the MRS Code of Conduct:

1. Researchers shall ensure that participation in their activities is based on voluntary informed consent.

2. Researchers shall be straightforward and honest in all their professional and business relationships.

3. Researchers shall be transparent as to the subject and purpose of data collection.

4. Researchers shall respect the confidentiality of information collected in their professional activities.

5. Researchers shall respect the rights and well-being of all individuals.

6. Researchers shall ensure that participants are not harmed or adversely affected by their professional activities.

7. Researchers shall balance the needs of individuals, clients, and their professional activities.

8. Researchers shall exercise independent professional judgement in the design, conduct and reporting of their professional activities.

9. Researchers shall ensure that their professional activities are conducted by persons with appropriate training, qualifications and experience.

10. Researchers shall protect the reputation and integrity of the profession.
Definitions

Definitions from the MRS Code of Conduct:

Anonymisation:

Anonymisation is the process of removing, obscuring, aggregating or altering identifiers to prevent the likely identification using reasonable means of the individuals to whom the data originally related.

Child:

A child is a person under the age of 16.

Client:

Client includes any individual, organisation, department or division, including any belonging to the same organisation as the member, which is responsible for commissioning or applying the results from a research project.

Data Collection Process:

A data collection process is any process used to obtain information from or about participants. It includes, but is not limited to, interviews, questionnaires, discussion guides, and stimulus materials, as well as passive data collection.

Identity:

The identity of a participant includes, as well as their name and/or address, any other information which offers a reasonable likelihood that they can be identified by any person or organisation who has access to the information.

Informed consent:

Informed consent is a process by which a participant voluntarily confirms his or her willingness to take part in a particular project, after having been informed of all aspects of the project that are relevant to their decision to participate.

Member:

A Member is an individual who has been admitted to membership of MRS in one of the categories set out the MRS Articles of Association.
For the purposes of applying this Code, an organisation with MRS members that has signed the MRS Company Partner Service Quality Commitment that applies throughout the organisation shall be treated as a Member.

**Monitoring**

Monitoring is the supervising of activities (such as data collection) to ensure that they meet required objectives and performance targets.

**Participant:**

A participant is any individual or organisation from or about whom data are collected.

**Research:**

Research is the collection, use, or analysis of information about individuals or organisations intended to establish facts, acquire knowledge or reach conclusions.

**Responsible Adult:**

A responsible adult is an individual who has personal accountability for the well-being of a child, for example a parent, guardian, teacher, nanny or grandparent.

**Definitions from the Data Protection Act 1998 used in the MRS Code of Conduct**

**Personal Data**

Data which relate to a living individual who can be identified-

(a) from those data, or

(b) from those data and other information which is in the possession of, or is likely to come into the possession of, the data controller,

and includes any expression of opinion about the individual and any indication of the intentions of the data controller or any other person in respect of the individual.

**Data Subject**

An individual who is the subject of personal data.

**Data Controller**
A person who (either alone or jointly or in common with other persons) determines the purposes for which and the manner in which any personal data are, or are to be, processed.

**Data Processor**

Any person (other than an employee of the data controller) who processes the data on behalf of the data controller.

**Processing**

Obtaining, recording or holding the information or data or carrying out any operation or set of operations on the information or data, including—

(a) organisation, adaptation or alteration of the information or data,

(b) retrieval, consultation or use of the information or data,

(c) disclosure of the information or data by transmission, dissemination or otherwise making available, or

(d) alignment, combination, blocking, erasure or destruction of the information or data.

**Third party**

Any person other than—

(a) the data subject,

(b) the data controller, or

(c) any data processor or other person authorised to process data for the data controller or processor.
1: The Interviewer and the Participant – gaining and maintaining co-operation

**The Rules**

10. Members must not disclose the identity of clients or any confidential information about clients without the client’s permission, unless there is a legal obligation to do so.

11. Where files of identifiable individuals are used, e.g. client databases, members must ensure that the source of the personal data is revealed at an appropriate point, if requested by participants. This overrides the right to client anonymity.

18. Members must ensure that participants are provided with sufficient information to allow informed consent to be given. This includes:

   a. the name of the organisation or individual responsible for data collection;

   b. the general subject of the data collection;

   c. the purpose of the data collection;

   d. whether the data collection is to be recorded and/or observed;

   e. who is likely to have access to live or recorded information;

   f. the likely length in minutes of the data collection, if asked;

   g. any costs likely to be incurred by the participant; and

   h. an assurance that the activity is being conducted in accordance with the MRS Code of Conduct.

   *Comment: the organisation or individual responsible for data collection is the data controller as defined under the Data Protection Act 1998.*

33. Members must take reasonable steps to ensure all of the following:

   a. that data collection processes are fit for purpose and clients have been advised accordingly;
b. that the design and content of data collection processes are appropriate for the audience being researched;

c. that participants are able to provide information in a way that reflects the view they want to express, including don’t know/ prefer not to say where appropriate;

d. that participants are not led towards a particular point of view;

e. that responses are capable of being interpreted in an unambiguous way; and

f. that personal data collected are relevant and not excessive.

34. Members must ensure that participants are informed about any recording, monitoring or observation at recruitment and at the beginning of a data collection process.

Comment: This does not include monitoring (listening to but not recording) telephone interviews for the purpose of quality control where interviewers have been informed that such monitoring takes place.

35. Members must ensure that participants are not misled when being asked to take part in a project.

36. Members must ensure that a participant’s right to withdraw from a project at any stage is respected.

Guidance

1. A good questionnaire should engage the participant from the start. Technically, the quality of data depends entirely on participant engagement and interest in the subject matter. Boredom, irrelevance, and questioning that are outside their frame of reference are likely to lead to poor data.

2. In addition, justice should be done to the knowledge and agenda of the participant. Researchers should consider how the participant would feel not only at the start of the interview, but during it, and when they have finished. How will they react – and will they be willing to help - the next time they are asked to take part in a research project; will they feel it had been an enjoyable experience; will they feel they had the opportunity to have their say about the topic; would they recommend taking part in research to others as a result of their experience?
3. There are occasions on which the research design requires the precise purpose of the research to be disguised. In such cases, the introduction need not explain the precise objectives of the research, but must honestly explain the broad subject matter. So, for example, in dealing with a subject such as cinema going, it may be legitimate to communicate the subject matter as being about “leisure activities”. It is important however to ensure that the description of the subject matter doesn’t mislead participants, in breach of B17.

4. The subject of the interview should be described as precisely as possible and be compatible with the objectives of the study. If any sensitive data is to be collected this must be made clear to the participant at the time when co-operation is sought. Researchers should apply their professional judgment in determining what could be sensitive in any given project. This includes, but is not limited to, subjects defined as “sensitive personal data” in the Data Protection Act 1998:

   a. racial or ethnic origin
   b. political opinions,
   c. religious beliefs or other beliefs of a similar nature,
   d. trade union membership
   e. physical or mental health or condition,
   f. sexual life,
   g. the commission or alleged commission of any offence, or
   h. any proceedings for any offence committed or alleged to have been committed, the disposal of such proceedings or the sentence of any court in such proceedings.

5. The objectives of a research project do not give the researcher a right to intrude on privacy or abandon normal respect for an individual’s values. This means that sensitive questioning/subjects must be particularly carefully handled.

6. In terms of sensitivity, topic areas can be divided into:

   - those topics that are judged sensitive to everyone, because of the nature of that subject;
• those topics that may be sensitive to a group for cultural, religious or political reasons; and

• those topics that may be sensitive to a particular individual, because of that individual’s past history, behaviour and attitudes.

7. In the case of the latter, researchers cannot necessarily take precautions in advance of the interviews, but can treat each case sensitively and individually, giving participants a genuine opportunity to withdraw. Any topic can potentially be sensitive to an individual. Participants who feel their privacy and personal sensitivities are not being recognised or respected may be less forthcoming. Equally, participants who feel they have not been treated with honesty and openness may feel patronised. In both cases, the nature of their responses may be affected. In addition, these participants may feel less willing to participate in further research.

8. In accordance with B18, All participants must be allowed to exercise their right to withdraw from the interview at any point. Further B14 provides that questionnaires must allow participants to express the views including “don’t know”, or “prefer not to say” where appropriate. In face-to-face, telephone or online interviews this means that the interview should generally be able to proceed without a pre-coded answer having to be selected.

9. Four major issues were identified by research undertaken by the Research Development Fund to impact negatively on both quality of data collected and participants’ attitude towards research. These are

• Excessively lengthy questionnaires, or a lack of honesty/transparency about the potential length of the interview

• Repetitive questioning

• Insufficient opportunity for participants to have their say

• Excessive classification section

10. Lengthy interviews can impact on response rates and hence representivity of the sample. Repetitive questions can negatively affect the participant’s willingness to be interviewed in the future and hence adversely affect future response rates.

11. Researchers should avoid long or repetitive interviews. This may involve a trade off against the “ideal world” objectives. Consider whether or not a question is required in order to meet the objectives, or whether it is “nice to know”.
12. Participants should feel that justice has been done to their knowledge and/or opinions within the confines of the questionnaire or subject matter. This may involve compromise on the part of the researcher.

13. It can be useful to think of the questionnaire as one element in a conversation. It should also be acknowledged that for the majority of participants some subjects (holidays or cars for example) are likely to be of more interest than others (household appliances or financial services for example). Researchers should take account of this and make particular efforts when dealing with more problematic topics.

14. A questionnaire designed for one interview methodology may need to be amended to be effective in another channel. However, researchers should be aware that may also lead to different answers across channels if adequate care is not taken.
2: Meeting the Objectives

The Rules

8. Members must take reasonable steps to design research to the specification and/or quality standards agreed with the client.

9. Members must take reasonable steps to ensure that the rights and responsibilities of themselves, clients, and sub-contractors are governed by a written contract and/or internal commissioning document.

Comment: The Data Protection Act 1998 requires contracts for the processing of personal data to be in writing.

33. Members must take reasonable steps to ensure all of the following:

a. that data collection processes are fit for purpose and clients have been advised accordingly;

b. that the design and content of data collection processes are appropriate for the audience being researched;

c. that participants are able to provide information in a way that reflects the view they want to express, including don't know/ prefer not to say where appropriate;

d. that participants are not led towards a particular point of view;

e. that responses are capable of being interpreted in an unambiguous way; and

f. that personal data collected are relevant and not excessive.

Guidance

1. Researchers should ensure that the objectives of a research project are fully translated into a series of questions, which will, as far as possible, obtain the required data.

2. It is the responsibility of all researchers involved in designing a questionnaire to ensure that the questions included are adequate to enable the study to meet its objectives. However where a researcher's advice and recommendation is not
accepted, the researcher cannot then be held responsible for any consequent failure to meet the objectives.

3. It is the responsibility of both researchers and clients to ensure that the questions asked meet the objectives of the study:
   - The researcher because they have a professional responsibility; and
   - The client because their commercial and professional interests are at stake.

4. The interests of research at large are thus served through the encouragement of:
   - The credibility of research in the commercial environment
   - Participant goodwill through a transparent and effective questionnaire.

5. Good practice dictates that questionnaires should be formally agreed to confirm that all parties are satisfied that the research objectives will be sufficiently met within the practicalities of getting the project done, paying particular attention to:
   - Errors of commission (objectives that are badly addressed)
   - Errors of omission (objectives that are not addressed either at all or in part).

6. Clients should obtain the guidance of researchers as to the feasibility of meeting defined research objectives:
   - Within an interview of a given length
   - Within participants’ ability to give the information required
   - Within ethical and technical best practice.
3: Structuring the questionnaire - its impact on interviewers and participants

The Rules

33. Members must take reasonable steps to ensure all of the following:

   a. that data collection processes are fit for purpose and clients have been advised accordingly;

   b. that the design and content of data collection processes are appropriate for the audience being researched;

   c. that participants are able to provide information in a way that reflects the view they want to express, including don’t know/ prefer not to say where appropriate;

   d. that participants are not led towards a particular point of view;

   e. that responses are capable of being interpreted in an unambiguous way; and

   f. that personal data collected are relevant and not excessive.

Guidance

1. Whenever possible, a questionnaire should follow some form of natural flow, reflecting a train of thought, a logical conversation, or an implied chronology of events, depending on the subject matter.

2. Good practice is to give high priority to user friendliness in administration, and a structure that is easy to follow for interviewers and participants.

3. Researchers should plan a questionnaire to make it logical from the participant’s point of view. This should involve outlining a structure and routing plan before beginning to write questions. The structure need not follow that of the objectives.

4. Consideration should be given to using blocks of sub-group questions and the construction of decision trees or flow charts to assist in structuring (and later routing) a questionnaire.

5. In general, order of questions will have an effect on the answers that are provided. This is generally controllable in an interview setting, or electronically, but when the
questionnaire is self-administered, it may not be possible to control the order in which the questions are read and answered.

6. **Avoidance of non-sequiturs** (i.e. statements having little or no relevance to what preceded them) is recommended. Where they are unavoidable (e.g. in omnibus style questionnaires) appropriate introductory phrases should be used.

7. To prevent a questionnaire being boring use a variety of different forms of questioning to break the routine within the questionnaire.

8. A theoretically optimal research design can result in a repetitive or boring interview. Researchers should, wherever possible, seek ways of engaging and maintaining the participant’s interest.

9. Question rotations between interviews, although good practice in themselves, are not a solution to this issue and sometimes have the effect of merely spreading unreliability across several questions, rather than concentrating it amongst a few.

10. The main principle is the rigour, integrity and quality of data. Neither participants nor interviewers should be confused or bored by any questionnaire.

11. Wherever possible, a questionnaire should be piloted before the survey to ensure it can be answered by potential participants as well as meeting the needs of the researcher/client (see Section 6)

12. The guiding rule can be expressed as “doing justice to how people would normally talk about things in everyday language”. 
4: Writing the Questionnaire

The Rules

33. Members must take reasonable steps to ensure all of the following:

a. that data collection processes are fit for purpose and clients have been advised accordingly;

b. that the design and content of data collection processes are appropriate for the audience being researched;

c. that participants are able to provide information in a way that reflects the view they want to express, including don’t know/ prefer not to say where appropriate;

d. that participants are not led towards a particular point of view;

e. that responses are capable of being interpreted in an unambiguous way; and

f. that personal data collected are relevant and not excessive.

Guidance

1. In summary, will the participants:

   • Understand the questions and /or the answer options?
   
   • Be willing to answer the questions?
   
   • Be able to answer the questions?

2. In practice every researcher knows that there is no right and wrong question or answer, and that, legitimately, different questions get different answers. The prescription for writing questions is therefore intrinsically linked to the interpretation of the results and data that will be obtained.

3. It is possible to frame questions or question sequences to support or dismiss any point of view, with varying degrees of subjectivity built in. This is particularly relevant to published research, and question wording is particularly crucial in opinion and advocacy research. The general public expects ‘objectivity’ in market research exercises even when this is not always the purpose.
4. Researchers should take particular care to ensure that participants are not lead to a particular answer. For example, any underlying facts to be communicated to the participant should be set out in a neutral manner. Positive phrases such as “the proposal is a good idea” in a question without a balancing negative phrase may lead the participant to a positive view of the proposal under consideration.

5. Sensitive subject areas should be explored through appropriately sensitive or even indirect questions. Sensitive subjects are most often placed towards the end of a questionnaire.

6. Participants should always be given the opportunity to decline to answer any question.

7. Any question that a participant is required to answer should be:
   - Within their frame of reference (in business-to-business research for example, can a single informant answer all the different questions?)
   - Relevant to them.

   If this is not achieved, participants will be forced to guess. In interpreting the responses, the researcher may not be aware of this.

8. Question concepts should be clear and understandable with as little ambiguity as possible. In general terms, a question should be interpretable in only one way.

9. Questions should be clear and phrased in language appropriate to the participant’s way of thinking/talking.

10. Only one question should be asked at a time - questions containing multiple concepts (e.g. What do you think about the colour and taste of the product?) rarely give sensible data.

11. The status of answers should be understood by researchers. Are we asking questions that can be answered:
   - Accurately (e.g. What was your age last birthday?)
   - Through memory (e.g. How many times in the last month have you done X . . .?)
   - Through a best choice of options, none of which may correspond precisely to the participant’s view / behaviour (e.g. Which of these three statements comes closest to describing your view . . .?)
• Through estimation, guesswork or even speculation.

12. Appropriate answer code options should be available that reflect the reality of the range of responses. Response codes should be:

• Clear and understandable, with as little ambiguity as possible;

• Phrased in language appropriate to the participant’s way of thinking/talking.

13. The questionnaire should be designed to indicate that a question was asked, even if it has not been answered. It should always be possible to record a “don’t know” response as this is often a perfectly valid answer, and a “not answered” response. The responses are distinct and should not normally be combined at the data collection stage. Classification questions can be seen as intrusive, and should be kept to a minimum.

14. Wherever appropriate space should be provided to allow the recording of comments of participants on the topic(s) not covered by the questionnaire. This achieves two objectives:

• It encourages participants to feel researchers are interested in and value their views;

• It might prove beneficial to the study in illustrating aspects of the topic(s) not adequately covered in the questionnaire.

15. Consider the appropriateness of question wording to the audience, particularly where it might be difficult to be answered by, or cause offence to, minority groups or people with special needs.

16. Where appropriate standard questions or questions used on previous research should be considered. Use of such questions gives comparability across studies and can enhance the value of the data to the client.
5: Questionnaire Instructions

**Guidance**

1. The instructions within a questionnaire are crucial to the reliability of the data returned and should be clear and unambiguous. Such instructions help both the interviewer (if used to gather the data) and the participant.

2. Each question in a questionnaire has four key routing instructions (whether overt or implied) associated with it:
   - Who should answer the question?
     For example: “Ask all” versus “Ask only those using product X at QY”.
   - How is the question to be asked?
     For example: “Read out” versus “Do not read out”.
   - How is the answer to be recorded?
     For example: “One code only” versus “code all answers mentioned”.
   - Where will they go next?
     For example: “Users ask QX” or “Non users skip to QY”.

3. The same design principles apply whether the questionnaire is administered on paper, by a computer-assisted method, or online.
   - In CAPI, CATI, online questionnaires or related computer assisted modes much of the routing will be embedded in the computer script and therefore need not be overt to the interviewer or participant.
   - Instructions for self-completion questionnaires should be clear and user friendly.

4. Neither interviewers nor participants should be held responsible for data collection errors arising from poor, or inadequately signposted, routing and/or instructions. Excessive complexity of routing is to be particularly discouraged. Consider including a question twice on the questionnaire in order to avoid complex routing.
6: Piloting

Guidance

1. On quantitative surveys, a small number of pilot (test) interviews should ideally be conducted to test the feasibility and appropriateness of the questionnaire. The main principle at stake is the responsibility of the researcher to collect high quality data in an objective manner, but there are also participant issues. It may be necessary to test whether the questionnaire is compatible with the rights of participants, particularly where sensitive issues or subjects are involved.

2. Where a client researcher is present at the pilot, the same rules apply as in other observed research.

3. Researchers should pilot questionnaires, not least to test the questionnaire for length, comprehensibility, and general good sense.

4. Researchers should satisfy themselves as to the workability of a questionnaire through an internal pilot, that is, pilot interviews with colleagues or others not involved in the research.

5. Researchers should conduct pilot interviews with a small selection of the target audience for any given research project.

6. Certain circumstances demand particularly rigorous piloting. For example:

   - When dealing with unfamiliar concepts – observation (watching pilot interviews), or qualitative pilots, can help in framing questions in the right way;

   - If there are areas in the questionnaire where there are no ready-made code lists, pilots can help generate initial codes for open ended questions;

   - Where it is obvious that the interview is potentially very complex or lengthy;

   - Where questions are potentially contentious;

   - When it is necessary to review and test alternative question strategies;

   - Where a questionnaire will be delivered on a variety of platforms, eg multiple web browsers, multiple operating systems or on mobile devices.
7: Dealing with the Results

The Rules

47. Members must ensure that findings disseminated by them are clearly and adequately supported by the data.

51. Members must provide clients with sufficient information to enable clients to assess the validity of results of projects carried out on their behalf.

52. Members must ensure that data include sufficient technical information to enable reasonable assessment of the validity of results.

53. Members must ensure that reports include sufficient information to enable reasonable assessment of the validity of results.

54. Members must ensure that outputs and presentations clearly distinguish between facts, opinion, and interpretation.

Guidance

1. Researchers have a responsibility to deal with results in a way that reflects the questions that were asked, rather than (mis)interpreting data in the abstract. This includes paying close retrospective attention to question phrasing, to whether the answers were prompted or unprompted, and so on.

2. Researchers also have a broader responsibility to deal with answers in a way that respects participants’ views. This will help retain the confidence of the general public and business community, ensuring future participant co-operation and the health and good image of the research profession.

It is impossible to divorce good practice in questionnaire design – in terms of ethical and technical responsibilities – from the interpretation of the resulting data. As part of the questionnaire design process, researchers should consider how they expect to analyse and report the results. This will aid in avoiding errors of both commission and omission.
8: Some references for help in questionnaire writing.


The Practice of Market Research, Chapter 9: Designing Questionnaires, Yvonne McGivern, FT Prentice Hall/MRS, 2009

ANNEX: Ten Tips for DIY Surveys

What?

Start with the end in mind. For each question you should be able to answer ‘what will I do when I have the answer’. How will it add insight and understanding? Be really strict on this, many questionnaires are too long because they fail to distinguish interesting from relevant.

Who?

Who do you want to collect the data from? Many DIY surveys canvas opinions from those in their own network, contact book or organisation. Their views may be representative of a wider population on some topics, but not all. Take care before assuming the results can be said to represent a wider group of people, particularly if you are taking decisions that will impact a broader group.

Set them up.

Today, it is common to be asked to complete a survey and many organisations ask customers for feedback from the last contact. The result is that such requests are all too easy to ignore. It helps to explain who you are, why you want their input and what you will do when you have the results. Also, tell people how long the survey will take. These things help achieve a higher response rate.

Who took part?

With online research it is now much easier to collect lots of responses. As well as the total number, take a good look at the profile of people who completed the survey. Do you have a good spread of participants, from different demographic groups and people who have roles with diverse experiences? Customers verses non customers or engaged verses disengaged participants can be expected to have different opinions.

When?

The timing of the DIY survey is important. People’s opinions change over time and on some topics can change rapidly. For example, when the subject concerns celebrity or music the shelf life of the data collected can be extremely limited. If possible allow a survey to run for some days to maximise response, but feel free to set an end-date.
How?

Be balanced and non-judgemental in how you phrase the question. You want to know what your audience truly think and not ‘lead the witness’. If you direct the question to favour one answer over another, the results are worthless and your audience may be frustrated at the bias.

Cover all the bases

People taking a DIY survey need to be able to answer your question with an answer choice that represents their view or position. Consider ALL the possible answers, not just the most common responses. It is good practice to offer ‘don’t know/don’t want to answer’ and, where it is difficult to predict the full range of answers, you might provide a category ‘other, please specify’ with a box to enter free text.

Use simple, straightforward language

Plain, unambiguous words and phrases are the safest route. The group from which you are seeking opinions will vary in their level of expertise, interest and commitment to the survey. If the question is complicated in its construction, the risk of misunderstanding is high. Before launching try the questions with a friend or colleague.

Ask open-ended questions

If your intention is to explore a subject, open questions like ‘why do you say that?’ help to reveal a range of opinions on a topic. While this is harder to translate into percentages, this approach will help to ensure you have the breadth of views people covered. Closed questions with multiple answers provide an alternative, but often require your audience to do some force-fitting of their perspective to your survey.

Share the results with care

The presentation of results as statistical totals implies rigour and accuracy. With any Research, care is needed to avoid over-interpretation of results. Note the sample definition, method and timing as well as the number of people completing the question. It is fine to be selective in highlighting results, but be careful not to distort the results. Good research is often described as a blend of art and science.