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Data Exchange Protocols (Version 6, October 2019)
1 Introduction

1.1 Purpose of this document

This document provides operational guidance to users of the Data Exchange across the full range of grant funded services, such as those associated with the Community Grants Hub and hosted by the Department of Social Services (DSS). The Data Exchange Protocols (the protocols) should be read in conjunction with the:

- Data Exchange Framework which outlines the principles and vision underpinning the Data Exchange
- organisation’s funding agreement
- task cards and e-Learning modules for users of the Data Exchange web-based portal
- Data Exchange technical specifications for all users submitting their data through system-to-system transfer or bulk upload from their own case management software.

The protocols are not intended to prescribe how organisations should run their business or collect data; they are intended to provide practical information for managers and front-line staff to help them integrate the Data Exchange data definitions and requirements into existing service and administrative practices.

The protocols are periodically updated to provide current and accurate guidance.

1.2 The Data Exchange Framework

The Data Exchange Framework represents the program performance reporting approach for client facing funding agreements. The Department of Social Services (DSS) progressively introduced standardised, prioritised, and collaborative reporting processes across grants programs from 1 July 2014. From 1 July 2016, the Data Exchange was expanded as part of the Commonwealth Government’s Community Grants Hub.

The requirements of the Data Exchange are divided into two parts: a small set of mandatory priority requirements that all organisations report, and an extended data set, known as the partnership approach. As part of the partnership approach, organisations are encouraged to collect the Standard Client/Community Outcome Reporting (SCORE) information for as many clients as practical.

This approach to reporting is streamlined, automated and includes a shift in focus of performance measurement from ‘outputs’ to more meaningful information about service delivery ‘outcomes’ through:

- **Streamlined reporting arrangements**—A standard client level data record (the priority requirements) applies across the broad suite of government funded client-based programs, replacing aggregate data reports, ‘smart forms’ and multiple IT reporting systems.

- **Free access to a web-based portal**—Organisations who do not have their own case management tools can access a free IT system. This helps record clients, service and outcomes data that meet funding agreement performance data requirements and allows organisations to confidentially manage their core client and case information.

- **Bulk uploading and system-to-system transfers**—The Data Exchange supports organisations who have a compatible case management software to transfer information directly from their own systems through bulk uploading and system-to-system transfers.

- **Promoting a partnership approach to reporting**—Organisations participating in the partnership approach to share client outcomes data with their funding agency in exchange for relevant reports. These reports are outcomes focused and include a rich set of added information to help inform service delivery using program performance, client survey and government data.

Go to the Data Exchange website for more information about the Data Exchange’s policy principles and program specific guidance.
2 Recording client level data

This section describes the important concepts and terminology associated with collecting and reporting client level data. It is important that managers and front line staff understand these concepts because they underpin the framework of the Data Exchange.

2.1 Client level data

Client level data refers to data collected and reported on each individual client rather than as summary (aggregate) data. The Data Exchange is designed to capture individual client level data. However, where collecting client data is not practical or possible, such as an activity involving a large group of people or a whole community, aggregate reporting is still accommodated by the system.

The main advantages of client level data are the:

- flexibility to analyse and report administrative data in multiple formats for different audiences, without burdening organisations with multiple data requests
- improved reliability of administrative data, as all organisations collect the same raw data records without the need to apply complex counting rules
- improved usefulness of administrative data, due to the use of a Statistical Linkage Key (SLK) allowing for the matching of de-identified data records across funded activities
- improved capacity for reporting data back to organisations, as the de-identified administrative data is held within a common data repository.

Client level data provided through the Data Exchange is de-identified so that no identifiable client information is used by an organisation’s funding agency.

Data Exchange staff work with organisations to ensure clear information is available to clients to affirm that only de-identified data is captured as part of program performance reporting and used for the purposes of policy development, grants program administration, research and evaluation.

Go to the Data Exchange website Information for clients about privacy and Section 4 of this document for more information about privacy. Program specific guidance on clients, support people and other client level data items is available on the Data Exchange website.

2.2 Who is a client?

When delivering program activities the term ‘client’ is used in many different ways, covering individuals, families, groups, other organisations and whole communities; as well as casework, participants, audiences and one-off contact. While this flexibility appropriately reflects the diverse strategies used to deliver services, we recognise that it may also create particular challenges for program performance reporting.

The Data Exchange uses a specific definition of ‘client’ to ensure comparable information is reported for the number of individuals that have received a service within a reporting period. This means that ‘apples with apples’ comparisons are possible within and across activities.

For the purposes of recording a ‘client’ record in the Data Exchange, a client is defined as:

An individual who receives a service as part of a funded activity that is expected to lead to a measureable outcome.

This definition includes a number of components that must be met in order to count a person as a client. These components are program and context specific and involve determining whether the individual in their own right is expected to achieve an outcome that is linked to a program specific objective.

Many different types of outcomes are achieved as part of service delivery. Outcomes are not limited to high-level, life changing events. Client outcomes can also be as simple as learning a new skill, receiving a service that is required, or gaining increased knowledge about other services that are appropriate and available. These outcomes should be recorded within the Data Exchange.
In cases where an organisation is funded to provide training, support and skills development of organisation’s staff, the staff member is considered a ‘client’ for the purposes of reporting in the Data Exchange.

2.3 Who is a support person?

At times, there may be other people present at a service who do not meet the definition of a client. This could include carers of clients, family members or children who attend to support the client. The support person is not expected to achieve a direct outcome through this service interaction and is not counted as a client.

There are no requirements to record the details of support people in the Data Exchange, however if an organisation wants, they can create an individual record for these people and record them as support people at the session level.

2.4 Services for children

A child is recorded as a client in the Data Exchange if the child directly received the service and meets the above definition of an individual client. When children are recorded within the Data Exchange the organisation should seek consent from a child’s parent or guardian, except in circumstances where the organisation considers that the child is a mature minor and is able to provide informed consent on their own.

2.5 Services for couples, families and households

The Data Exchange captures information about individual clients, however there are some funded activities where multiple individuals are assisted as part of the same ‘case’, ‘family’ or ‘group’. In these instances, a client record should be created for each individual client and grouped together using a ‘case’ record.

Table 1. Example of a client and support person

<table>
<thead>
<tr>
<th>Activity/Service Context</th>
<th>Who is the client?</th>
</tr>
</thead>
<tbody>
<tr>
<td>Helping a couple at risk of homelessness to manage their finances in order to prevent a future accommodation crisis.</td>
<td>Both people in the couple are considered clients, as they are both receiving the service, benefit from that service, and meet the definition of a ‘client’ as per the program activity guidelines. Two client records should be created and used within the Data Exchange.</td>
</tr>
<tr>
<td>A youth attends counselling and their parent/carer is in the room during the counselling session.</td>
<td>The youth is counted as the client as they have received the service and will achieve an outcome. The parent/carer is not recorded as a client as no measureable outcome is achieved on this occasion. The parent/carer could be recorded as a support person, however this is not mandatory. If the parent/carer attends a session and participates directly (achieving their own outcome) they would be recorded as clients.</td>
</tr>
</tbody>
</table>

2.6 Recording unidentified clients

Some organisations deliver services to groups in the community. These activities can include information sessions and public events. The purpose and delivery of this work differs across activities but can include early intervention and prevention, education, awareness raising and capacity building. Sessions can range in size and there may be identified clients (where a client record is created for each individual attending), and unidentified clients (where an aggregate attendance figure is recorded) or a mix of both.

Organisations are strongly encouraged to create individual client records for as many of their clients as possible. If it is impractical to collect information about individual participants, for example in community outreach activities where many members of the general public may participate, the aggregate number of unidentified clients is recorded.

Unidentified client numbers captured at the case level estimate the total number of unidentified clients who are anticipated to attend services. In contrast, the unidentified client attendance numbers captured at the
session level records how many unidentified clients actually attended that instance of service. This prevents double counting.

For example, if the unidentified client total at the case level is 100, and there are two sessions each with 100 unidentified clients; the total specified at the case level indicates that there may be up to 100 unique unidentified clients for each session, and not 200 in total.

In some instances, there may be a combination of unidentified clients and individual clients in the same case and sessions. For example, an information session may have a number of identified clients as well as a number of unknown new clients or members of the general public. Identified clients should always be recorded as attending, as well as the unidentified clients recorded using the unidentified client attendance field. This number is reported against both the case and session records.

Collecting unidentified clients demonstrates how many people attended an event, but does not include any further data (such as Aboriginal and Torres Strait Islander origin, culturally and linguistically diverse (CALD) background, age groups).

For organisations participating in the partnership approach, the outcomes achieved from group activities are recorded through the Community component of the Standard Client/Community Outcome Reporting (SCORE).

### Table 2. Example of when to create a community or group session

<table>
<thead>
<tr>
<th>Activity/Service Context</th>
<th>Can unidentified ‘group’ clients be recorded?</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mental health awareness event for carers and families in the community. These were individuals attending information and advisory workshops.</td>
<td>Yes. As anonymous members of the public, these 25 people are recorded as an aggregate count (unidentified clients) because it is not practical to record individual client details or measure individual client outcomes.</td>
</tr>
<tr>
<td>A group attends a public event in the community. These were individuals attending an event or help with parenting issues and to find relevant programs in the service.</td>
<td>Yes. Before the event, the service will create a case record to indicate the number of expected participants. After the event, the service will report the actual participants at the session level. Of these participants, some signed up for programs offered by the service and are expected to achieve a measurable outcome in their parenting skills. A client record is created for these participants. The other participants have no further relationship with the service and are recorded as unidentified clients.</td>
</tr>
</tbody>
</table>

### 2.7 Group session

When delivering program activities, a group session generally means a session that has three or more clients attending a session together. A group session can be made up of a family group, clients that are known to each other, or strangers. This definition does not count a support person or practitioner as a member of that group.
3  Linking client data to service delivery

3.1  What is a service?
The Data Exchange framework has a specific definition of a service based on service delivery concepts. These concepts ensure that an instance of service is consistently applied across varying funded activities and service delivery contexts that are reported in the Data Exchange. For the purposes of the Data Exchange, a service is defined as:

One or more individual instances or episodes of assistance (known as sessions) within a reporting period that are delivered within a distinct case.

The concept of a ‘case’ and ‘session’ are integral to the Data Exchange as they maintain a consistent way for organisations to record information about the different activities clients are accessing, how they are being delivered and the location from which they are being delivered. These concepts are discussed below and in further detail at Section 5 of this document.

Go to the Data Exchange website program specific guidance for more information on cases and sessions.

3.2  What is a case?
Cases act as containers, linking client and session data to location and program activity information. A case is defined as:

A method to capture one or more instances of service (known as sessions) received by a client or group of clients that is expected to lead to a distinct outcome. A case may contain between one and an unlimited number of sessions.

A case record helps understand what funded activity is being delivered, the location it is being delivered from, the reason clients came to the service and the number of clients receiving a service.

Each organisation can create cases in a format that best suits their needs. For users of the web-based portal, cases facilitate navigation and hold clients and sessions together.

- A case can operate over multiple reporting periods, for instance if a client returns to receive the same service.
- Depending on the nature of the service, a case can contain an individual, a couple, a family, or an unrelated group of individuals, such as a regular weekly or monthly group meeting.
- If a client attends a number of different funded activities, each of these is treated as a separate case.
- If a client receives the same services from a number of different locations (known as outlets) managed under the same program activity, each of these is treated as a separate case.
- To report a case, details are recorded about the activity, the location (or outlet) where the service occurred, and the client who will receive the service associated with that case record.

3.3  What is a session?
In the Data Exchange, a session is defined as:

An individual instance or episode of service, stored within a case which is ‘related’ to other sessions (when/if they occur).

A session record includes the date the service occurred, the kind of service the client(s) received (known as service type) and which of the clients associated to the case were present. For organisations participating in the partnership approach, client pathways information (referrals out) is recorded at a session level. More information about the extended data set is found in Sections 6 and 7 of this document.

For two specific service types (advocacy/support and record searches), an instance of service is recorded even if the client is not physically present or is only present on the telephone. This is due to the nature of the
service provided, and only applies if a substantive effort was put into providing the service and the client is directly benefiting from the service.

3.4 Counting rules for clients, cases and sessions

A client is counted against a reporting period if the client was recorded as attending at least one session within that reporting period.

A case is counted against a reporting period if at least one session is recorded under the case within that reporting period.

A session is counted against a reporting period if the date of the session fell within the reporting period and at least one client is attached.

3.5 What is an outlet?

For the purposes of the Data Exchange, an outlet is defined as:

The physical location from where a service is primarily being delivered.

- The organisation identifies the program activities each outlet delivers.
- Each outlet can have different staff, service information, program activities, and contact details.
- Where the service is mobile in nature, the outlet used should be the nearest administrative premises where staff are based, and where they are likely to be travelling from to deliver the service.
- Creating multiple outlets for services delivered from the same address must be avoided.
- Post office boxes cannot be used in place of a physical location.
- An outlet should never be created for a client’s residential address, if a service is delivered in a client’s home, or a sensitive/protected location such as a refuge.
- In the instance of service delivery at a residential address, the outlet should reflect where staff are based or travelling from. This information is captured with the session details under the service setting field.
- In the instance of service delivery at a protected address or refuge, the outlet can use an address of a non-identifiable public place nearby, such as a post office, police station or shopping centre.

3.6 Delivery partnerships and consortium arrangements

Organisations make different choices when it comes to setting up their delivery partners and outlets. As these decisions will affect who can enter, view and report data in the Data Exchange, set up needs to happen in agreement between the two parties, e.g. lead/facilitating and delivery partners. Particular attention needs to be paid to the naming of outlets, outlet addresses, the visibility of data and the protection of client privacy and personal information.

Go to the Data Exchange website training resources for guidance on partnerships and consortium arrangements.

3.6.1 Funding received from multiple sources

The Data Exchange Framework is intended to capture client outcomes from services funded through programs in scope for the Data Exchange. Where an organisation receives funding from multiple sources to help the delivery of an activity, the following guidance should be considered:

(a) Where organisations deliver activities that are funded through multiple sources, the data reported should reflect the clients interaction with in-scope programs over the whole funding or reporting period (as opposed to only seeing a snapshot over a shorter period).
(b) If Data Exchange related funding is only provided for a specific aspect of the service offering, such as in a certain location or to a specific client group, it is acceptable to only report on those clients or that specific outlet.

(c) When an organisation distributes a voucher or other specific items (such as Energy Accounts Payment Assistance (EAPA) vouchers) and these items are acquitted in relation to the agency that provided the item and this agency is not reporting through the Data Exchange, this is not recorded in the Data Exchange.
4 Protecting a client’s personal information

The Data Exchange Framework was designed to ensure a client’s personal information is protected through stringent protocols that comply with the requirements of the Privacy Act 1988, including the Australian Privacy Principles (APPs).

Under the Privacy Act, personal information is information or an opinion about an identified individual, or an individual who is reasonably identifiable:

(a) whether the information or opinion is true or not; and
(b) whether the information or opinion is recorded in a material form or not.

Information that is stored by DSS on the Data Exchange is ‘personal information’ if it relates to an identified individual, or an individual who is reasonably identifiable, and this information says something about them. For example, an identified individual on the Data Exchange includes data with the client’s name and their reasons for seeking a service from an organisation.

4.1 Data Exchange privacy protocols

The Data Exchange has been designed to accommodate the different ways organisations collect and report client-level data. Some organisations choose to use the Data Exchange to collect and store personal information, whereas other organisations do not.

Regardless of the upload method used (web-based portal, bulk upload or system-to-system transfer), organisations must apply the Data Exchange consent and notification arrangements if they intend to store personal information in the Data Exchange.

Where an organisation stores personal information in the Data Exchange, only the organisation can access the personal information stored on this DSS hosted information system. Strict IT security protocols prevent DSS staff from accessing personal information in this system for any purpose other than confirming that the privacy protocols are working correctly.

Funding agencies use de-identified data from the Data Exchange for program management, policy development, research and evaluation activities for government. DSS applies best practice data de-identification and aggregation methods when producing reports and information for these purposes, to ensure that a client cannot be identified or re-identified by other government departments or organisations.

Go to the Privacy Impact Assessment (PIA) on the Data Exchange website to find out more about the PIA assessment conducted by the Australian Government Solicitor which examined the Data Exchange’s compliance with the Privacy Act.

4.2 Organisations’ obligations when storing personal information in the Data Exchange

Organisations using the Data Exchange to collect and store personal information are considered to be using the Data Exchange as a “client records system” or “client management system”. This means they need to see the client’s personal information in the Data Exchange for the purpose of managing their client records and the services being delivered to clients.

When reporting client-level records and using the Data Exchange as a client records system, the personal information that organisations submit to the Data Exchange (either through the web-based portal, bulk uploading of files or a system-to-system transfer) is stored by DSS on the Data Exchange. This includes the client’s name/pseudonym and address information (line 1 and 2).

Organisations who use the Data Exchange as a client records system must adhere to the notification and consent requirements identified below, which ensure that DSS complies with its obligations under the Privacy Act and the APPs.

To satisfy the notification requirements organisations must include the DSS standard notification (paragraph 4.2.1 below) on their registration forms. If organisations do not wish to use these words on their registration
forms, organisations are required to notify the client of the matters outlined in APP 5.2, or ensure that the client is aware of those matters.

In either case, organisations are required to provide DSS’s standard notification (or an alternative notification on privacy) before the time that the client’s personal information is entered on the Data Exchange or, if that is not practicable, as soon as practicable after the client’s personal information is entered on the Data Exchange.

4.2.1 Standard notification on privacy

Organisations using the Data Exchange as a client records system must include the ‘DSS standard notification’ on their registration forms to enable DSS to store a client’s personal information on the Data Exchange. The DSS standard notification is outlined below:

“The information that we collect from you on this form includes your personal information. Your personal information is protected by law, including by the Commonwealth Privacy Act

The client management system that we are using is an IT system called the ‘Data Exchange’. This system is hosted by the Australian Government Department of Social Services (DSS). Your personal information that is stored by DSS on the Data Exchange will only be disclosed to us for the purposes of managing your case. You are not required to provide your personal information to DSS. If you do not consent to the collection of your personal information, this will not affect the services provided to you. If you provide your personal information to DSS, you can ask for this information to be removed by DSS at any time.

DSS de-identifies and aggregates data in the Data Exchange to produce information for policy development, grants program administration, and research and evaluation purposes. This includes producing reports for sharing with organisations. This information will not include information that identifies you, or information that can be used to re-identify you, in any way.

You can find more information about the way DSS will manage your personal information in DSS’s privacy policy, which DSS has published on its website. This policy contains information about how you may access the personal information about you that is stored on the Data Exchange and seek correction of that information. This policy also includes the circumstances in which DSS may disclose personal information to overseas recipients, as well as information about how you may complain about a breach of the Australian Privacy Principles by DSS, and how DSS will deal with your complaint.”

4.2.2 Alternative notification on privacy

If organisations do not wish to include the standard notification on their registration forms, they may design and use their own forms to collect and store personal information when using the Data Exchange as a client records system. If organisations wish to follow this approach, they are required to notify the client or otherwise ensure that the client is aware of the following matters (as outlined in APP 5.2):

(a) the Data Exchange is an IT system that is hosted by DSS
(b) the organisation is using the Data Exchange for client management purposes, and the client’s personal information is stored on the Data Exchange for this purpose only
(c) the client’s personal information which is stored by DSS on the Data Exchange, is only visible to the organisation that collected the information for the purposes of managing the client’s case
(d) DSS de-identifies and aggregates personal information that is stored on the Data Exchange to produce information for policy development, grants program administration, research and evaluation purposes, and this will not include information that identifies the client, or re-identifies the client, in any way
(e) DSS’s privacy policy is published on its website. The website contains information about how the client may access or correct the personal information that is stored on the Data Exchange; complain about a breach of the APPs by DSS, and how DSS will deal with the client’s complaint. The privacy
policy also contains information about the circumstances in which DSS may disclose personal
information to overseas recipients
(f) the consequences if personal information is not collected from the client.

This notification is necessary to enable the client’s personal information to be stored on the Data Exchange
by DSS in compliance with the Privacy Act. DSS will not review, approve or store an organisation’s
registration forms.

4.2.3 Consent to collect and store personal information

In addition to providing the required notification to a client, an organisation using the Data Exchange as a
client records system will need to:

- obtain the express consent of a client, on behalf of DSS, for DSS to collect the client’s personal
  information from the organisation and store it on the Data Exchange
- record that consent in the Data Exchange IT system
- inform the client that they may withdraw this consent at any time.

DSS will not store the client’s name or pseudonym, and/or street-level address on the Data Exchange unless
this consent is obtained.

To meet DSS’s obligations under the Privacy Act, the required consent is to be recorded in the Data
Exchange. For organisations using the web-based portal, consent is recorded using the tick box provided
when creating or editing a client. When reporting client-level records through bulk uploading of files or a
system-to-system transfer, this consent is to be recorded in these files.

Organisations are required to record that a client has consented in the Data Exchange, however they are not
required to provide copies of the client’s consent to DSS. Organisations should determine their own record
keeping procedures in relation to client consent.

If a client withdraws consent for DSS to store his or her personal information on the Data Exchange,
an organisation must record the client’s decision in the Data Exchange. Organisations using the web-based
portal will need to ‘un-tick’ the consent for personal information box. Organisations that report using system-
to-system transfers or bulk uploads will need to update the client level record in the Data Exchange
consistent with a client’s decision to withdraw their consent.

When an organisation has recorded a client’s decision in the Data Exchange, DSS will process the
withdrawal of consent by removing the record of the client’s name or pseudonym, and/or street-level address
from the Data Exchange.

4.2.4 Obtaining consent from children and individuals with compromised capacity

When obtaining consent from a client who is a child, it is best practice to seek consent from the child’s parent
or guardian, except in circumstances where the organisation considers that the child has sufficient
understanding and maturity to understand what is being proposed. As a general rule, the organisation may
presume that an individual aged 15 or over has capacity to consent, unless there is something to suggest
otherwise. An individual aged under 15 is presumed not to have capacity to consent.

When obtaining consent from a client whose capacity to consent may be compromised (e.g. a client with a
physical or mental disability), it might be appropriate to implement special practices. For example, the
organisation should consider who can provide consent on the client’s behalf. Options include a guardian,
someone with an enduring power of attorney, a person recognised by other relevant laws (e.g. a ‘person
responsible’ under the Guardianship Act 1987 (NSW)) or a person nominated in writing by the client while
they were capable of giving consent.
4.3 Organisations' obligations when not storing personal information in the Data Exchange

The consent and notification arrangements identified in section 4.2 do not apply to organisations if they choose not to store personal information in the Data Exchange.

For this purpose organisations will indicate that client consent has not been provided when reporting through a system-to-system transfer or bulk upload. The information that is stored in a client level unit record will not include the client's name, pseudonym or street-level address.

4.4 Consent for follow up research

Funding agencies and third parties (such as universities) are interested in commissioning future research to better understand client needs and find opportunities to improve service delivery. Obtaining client consent to participate in research will create an indicative pool of willing participants for future research projects.

Organisations must ask clients if they consent to participate in future client research. This consent forms part of the priority requirements. Future research will vary depending on the nature of the planned evaluation, however the basic steps are:

- Any research conducted will be approved by a recognised ethics committee.
- Researchers will communicate with organisations before any research activities start.
- Organisations and researchers commit to clear and simple communications to help clients understand why research is important and what it means to participate.
- Clients can withdraw their consent at any time.

4.5 Organisational privacy considerations

Once a client record is created it is visible to all Data Exchange users within the organisation. Organisations must address any potential privacy issues through their own internal business processes.

Organisations should never provide a client’s personal information to DSS via telephone or email communications, for example when contacting the Data Exchange Helpdesk.

4.6 De-identified data

The Data Exchange protects client privacy by applying best practice data de-identification and aggregation methods, including the use of statistical linkage keys for data matching.

4.7 Unique client identifiers

A unique client identifier is used to ensure client records are matched in the Data Exchange.

Two data items (date of birth and gender) are included in the priority requirements to help identify individual clients without disclosing personal information. Program performance data provided by organisations through the Data Exchange is de-identified and encrypted so that no personal client information is accessed by DSS or any other agency.

4.7.1 Statistical Linkage Key (SLK)

The Data Exchange de-identifies client data using Statistical Linkage Keys (SLK), an algorithm developed by the Australian Institute of Health and Welfare (AIHW). The SLK is a method that allows DSS to link clients with the service they received through a unique key, enabling two or more records belonging to the same individual to be linked. The Data Exchange uses SLK 581 as a unique client identifier.

The SLK is an algorithm that creates a code consisting of the second, third and fifth characters of a person’s family name, the second and third letters of the person’s given name, the day, month and year of the person’s date of birth and the gender of that person. For example, John Smith, a male born on 14th February 1971 has an SLK of: MIHOH140219711
A client’s SLK is not visible to organisations in the Data Exchange. The SLK is only visible to a highly restricted number of DSS employees who perform database administration or data analytics duties to ensure the IT system behind the Data Exchange remains functional and to support data sharing and reporting initiatives.

For organisations using the Data Exchange web-based portal, the SLK is automatically generated within the system. For organisations using bulk uploads or system-to-system transfers to upload data to the Data Exchange, the SLK can be incorporated into their own client management information system using the AIHW’s algorithm (SLK 581). This SLK must not include shortened versions of a client’s name, nick name or any variations of their full name, or any other information that could identify a client.

Go to the Data Exchange Web Services technical specifications for help configuring systems to push the SLK across to the Data Exchange.

4.7.2 Client ID

Each client record in the Data Exchange includes a client ID that must remain unique to that client in all circumstances. The client ID is different to the SLK that is described above. The client ID is a technical data item used to organise client records within the Data Exchange and does not include any personal information. The client ID is made up of a series of alpha numeric text, either inputted by the organisation or created by the Data Exchange web-based portal.

Organisations using the Data Exchange web-based portal will have the option of entering their own client ID (an ID used internally by the organisation to administratively manage clients) or leaving the field blank. The system will generate a client ID that is used by organisations to search for their record at a later time.

Where an organisation chooses to enter their own client ID, this should be alphanumeric or numeric text only. This Client ID must not include shortened versions of a client’s name, nick name or any variations of their full name, or any other information that could identify a client, under any circumstances.

For organisations using their own client management system and uploading their data to the Data Exchange through bulk uploads or system-to-system transfers, the client ID becomes a mandatory field that is pushed across from their own system and used to cross reference the record between the two systems in future interactions.

Go to the Data Exchange website for technical specifications.
5 Collecting the priority requirements

The priority requirements are a small set of mandatory data items. These data items capture the demographics of clients accessing program activities, how often clients are attending, where they are attending and what program activities they are attending.

In summary, the priority requirements reflect the collection of information about client details, case and session details, and client consent to participate in follow-up research.

This section presents practical information about each of these concepts to support managers and frontline staff to consistently and accurately collect the required data.

Go to the Data Exchange website for more information on configuring systems and service delivery information such as how to use cases, sessions and service types. The data values are listed in Section 11 of this document.

5.1 Client level data

The priority requirements capture client details and demographic characteristics. This provides an understanding of each client’s pathways over time, on a de-identified basis.

A client record only needs to be created once. It can then be maintained, updated and edited at any time.

Client level data is reported for all individuals who receive a service as part of a funded activity, in line with the definition outlined in Section 2 and 3 of this document. These records are the basic ‘building blocks’ of the Data Exchange Framework and are used to answer standard questions such as:

- how many clients were assisted?
- how many clients were assisted in previous reporting periods?
- how many clients received assistance under different funded activities?
- how many clients received assistance from a funded activity delivered by a different organisation?
- how many clients receiving assistance were from vulnerable target population groups?

Answers to these questions will help tell the broader story about the outcomes being achieved, by providing an understanding of who these outcomes are being achieved for and when.

5.1.1 Collecting client given and family names

Collecting a client’s name would typically occur the first time that a client accesses any funded activity from an organisation, either in a registration form or an intake interview. Organisations are free to gather this information in accordance with their standard practices.

A client’s given name and family name are recorded because they form part of the SLK used to uniquely identify clients without disclosing personal information. Given name is typically a client’s first name, but it may include one or more middle names. Ideally, the given name should be recorded exactly as it is on key identification documents such as a passport or driver’s license.

Family name is typically the client’s last name, or surname, and ideally should be recorded exactly as it is spelled on key identification documents.

Where clients are known by more than one name, or prefer to be called by a particular name, for example Joe rather than Joseph, their given and family names should reflect the name the client offers.

Where a client does not have identification documentation or chooses not to disclose their identifying information, the organisation should record the given and family name that is most commonly used or preferred to be used by the client.

Where a client does not wish to disclose their ‘real’ name, the organisation should indicate that a pseudonym is being used and record a pseudonym that ideally is used again if the client returns for other services. Where a client only has one name, this would be entered as their first and last name.
5.1.2 Date of birth

A client’s date of birth is recorded for two reasons: it forms part of the SLK and provides a direct means of calculating the client’s age.

Age groups demonstrate part of the standard demographic profile for clients required by many government programs and is of particular importance to programs that target age-specific cohorts.

Where a client does not know their date of birth or does not wish to disclose it, it is acceptable for an estimated date of birth to be used. An estimated date of birth indicator is in the Data Exchange and should be used to flag when this occurs. For example, if a client thinks they are approximately 30 years old (and it is 2019), the estimated date of birth indicator is flagged and the year of birth is recorded as 1989.

5.1.3 Gender

A client’s gender is recorded because it forms part of the SLK and is recorded based upon how the client self-identifies. Please note that gender is different to sexuality and sexual orientation which are not recorded in the Data Exchange.

The Data Exchange uses standard data definitions for gender developed by the Australian Institute of Health and Welfare (AIHW), with four options:

- Male
- Female
- Intersex/indeterminate
- Not stated or inadequately described

The ‘Intersex/indeterminate’ response is used where a client does not identify as male or female. If a client chooses not to disclose their gender, it is acceptable to record ‘Not stated/inadequately described’.

5.1.4 Residential address

Information about where clients live can assist with understanding if services are located in the right area.

A client’s residential address can also be compared to an outlet address to understand how far the client may be travelling to access a service, or how far staff may be travelling to deliver a service to a client.

A client’s residential address can also be linked to other useful information to help understand a client’s circumstances, such as the Socio-Economic Indexes for Areas (SEIFA) rankings and the Australian Bureau of Statistics (ABS) community profiles.

Within the Data Exchange there is the capacity to record a full residential address for each client. At a minimum, a client’s state, suburb and postcode are considered part of the priority requirements and must be recorded to create the client record.

The identity of clients providing their full residential address is protected by converting the data to the Australian Statistical Geography Standard. This means that a geography code is recorded in place of the client’s address, which de-identifies the record.

In exceptional circumstances, it may not be appropriate to record the client’s full residential address, such as where the client is experiencing domestic violence and does not wish to provide even their suburb, state and postcode due to fears for their personal safety. In these circumstances, the service outlet suburb, state and postcode should be recorded instead.

5.1.5 Recording a homeless client’s residential address

If a client is homeless or of no fixed address, the client or organisation can determine the most appropriate address to be recorded. This may be the suburb, state and postcode of where the client usually spends the night, or suburb, state and postcode of the outlet where the client is seeking assistance. A flag to indicate the client is currently homeless is in the extended demographics section of the Data Exchange.
5.1.6 Indigenous status

A client’s Indigenous status is recorded to provide an important understanding of whether clients who identify as Aboriginal or Torres Strait Islander origin are accessing services. Under standard data collection definitions used by the AIHW, five options are available to record a client’s Indigenous status.

Indigenous status is part of the standard demographic profile for clients of many government programs and is of particular importance in ensuring Indigenous people and communities have appropriate access to funded services.

Where a client chooses not to disclose their Indigenous status, it is acceptable to record ‘Not stated/Inadequately described’.

5.1.7 Cultural and Linguistic Diversity (CALD)

A client’s CALD background is recorded to provide an important understanding of whether CALD clients are accessing services. Under standard data collection definitions used by the AIHW, two questions are used to record a client’s CALD status:

(a) Country of birth
   - Record the country of birth indicated by the client
   - A list of values is based on the Australian Bureau of Statistics Standard Australian Classification of Countries (SACC), 2016

(b) Main language spoken at home
   - Record the main language spoken at home indicated by the client.
   - A list of values is based on the Australian Bureau of Statistics Australian Standard Classification of Languages (ASCL), 2016

More detailed information about a client’s CALD background such as ancestry is collected in the extended demographics section of the Data Exchange.

CALD status is part of the standard demographic profile for clients of many government programs and is of particular importance to ensure CALD clients and communities have appropriate access to funded services.

This information can also be beneficial for organisations in determining whether the engagement of translating services or bilingual staff may assist in better service delivery for their clients.

Where a client chooses not to disclose their CALD status, it is acceptable to record ‘Not stated/Inadequately described’.

5.1.8 Disability, impairment or condition

Clients are asked to self-identify whether they have a disability, impairment or condition because it is important for organisations and funding agencies to know whether clients with disability are accessing services.

Under standard data collection definitions used by the AIHW, disability is recorded in groupings that most clearly express the experience of disability by a person. Disability groupings constitute a broad categorisation of disabilities in terms of the underlying health condition, impairment, activity limitations, participation restrictions, environmental factors and support needs. Categories in the Data Exchange include:

- **Intellectual/learning**: associated with impairment of intellectual functions which limit a range of daily activities and restrict participation in a range of life areas, for example, but not limited to; dyscalculia, dysgraphia, dyslexia.

- **Psychiatric**: associated with clinically recognisable symptoms and behaviour patterns frequently associated with distress that may impair personal functioning in normal social activity, for example, but not limited to; Asperger syndrome, attention deficit hyperactivity disorder, autism, behavioural disorders, bipolar, depression, eating disorders, epilepsy, manias, phobias, schizophrenia, somnias.
- **Sensory/speech:** including vision disability (blindness, vision impairment); hearing disability (deafness, hearing impairment that cause severe restrictions in communication); deaf-blind (dual sensory impairments causing severe restrictions in communication); speech disability (speech loss, impairment which causes severe restrictions in communication).

- **Physical/diverse:** associated with the presence of an impairment, which may have diverse effects within and among individuals, including effects on physical activities such as mobility. This grouping includes physical disability, for example; paraplegia, quadriplegia, muscular dystrophy, motor neurone disease, neuromuscular disorders, cerebral palsy, absence or deformities of limbs, acquired brain injury, neurological disability (including epilepsy, dementias, multiple sclerosis and Parkinson disease).

- **None:** no disability, or no disability, impairment or condition are identified by the client.

- **Not stated/inadequately described.**

When recording data about disability, impairments or conditions clients should self-identify, and can identify with more than one group, for example physical/diverse and intellectual/learning.

Data about disability status is part of the standard demographic profile for clients of many government programs and is of particular importance to ensure people with a disability have appropriate access to funded services. Where a client chooses not to disclose if they have a disability, impairment or condition, it is acceptable to record ‘Not stated/Inadequately described’.

### 5.2 Service delivery information

The concept of a case and session are integral to the Data Exchange as they maintain a consistent way to link a client with instances of service and to help tell the ‘story’ about outcomes achieved for clients.

Once a client record is created in the Data Exchange, it must be linked to the program and activities the client is participating in. This is captured using the case and session records. A case is the first step in recording service delivery information within the Data Exchange.

#### 5.2.1 Case details

The second tier of the priority requirements is a case record, which includes a case ID, program activity, and outlet information. A case record is only created once for each unique case and is used over multiple reporting periods. Each case record includes:

- **Case ID:** an alphanumerical code or title that uniquely identifies the case, and which is named in a way that is meaningful to the user. The case ID business rules are the same as those for the client ID: the case ID must be unique within the organisation and not include any identifiable information, such as a client’s name or their Centrelink Customer Reference Number. Users of the Data Exchange web-based portal may leave the field blank, in which a case ID is automatically generated (numeric only). The field is mandatory for those uploading data through the bulk upload or system-to-system methods.

- **Program activity:** the funded activity that the case is being delivered under.

- **Outlet:** the location where the case is primarily being delivered. A case cannot have more than one outlet.

- **One or more client records:** links one or more clients to a case (or in limited circumstances an aggregate number of unidentified clients).

The number of case records an organisation creates will depend on the type of funded activity(ies) they deliver and the way these services are delivered. For example, if providing counselling to couples or families it would make sense to create a case for each couple/family. This would allow a user to see and reflect on the family composition of each couple/family, easily navigate the portal for efficient data entry, and potentially count the total number of cases as the number of couples/families accessing services.
In contrast, for organisations delivering activity-based services, it may be better suited to create a case for each of the locally run activities delivered in the community, such as a breakfast club or education course.

For organisations using their own client management systems, the concept of a case is a node that allows all three tiers of the Data Exchange data (clients, cases and sessions) to be effectively uploaded.

5.2.2 Session details

The third tier of the priority requirements is a session record. A session record captures the types of services being delivered under the relevant case, which clients attended, and the dates of service. Sessions also indicate that a case was active within a reporting period. Each session record consists of:

- **Session ID**: a numeric code or title that identifies a particular instance/episode of service. The session ID must be unique within the case and cannot include identifiable client information. Users of the Data Exchange web-based portal may leave the field blank and a session ID is automatically generated (numeric only). The field is mandatory for those uploading data through the bulk upload or system-to-system methods.
- **Session date**: the date the instance/episode of service occurred.
- **Service type**: the main focus for the session delivered. If a session covered multiple service types the most relevant one should be chosen, either based on the majority of time spent or the main way an outcome was achieved.
- **Client attendance**: recorded for each client that was present at the session.
- **Unidentified client attendance**: the aggregate number of clients who attended a session. This should be limited to large groups where the collection of client level information is not feasible. Unidentified client attendance at a session must be less than or equal to the number of unidentified clients against the case.

When recording a session, organisations should select the service type which best reflects the nature of service delivery in that particular session. Different service types are associated with different funded activities. Within the Data Exchange web-based portal only the relevant service types are available for a user to choose.

For organisations using their own client management systems, sessions are a node that complete all three tiers of Data Exchange data (clients, cases and sessions) being effectively uploaded.

5.3 Program specific mandatory fields

The Data Exchange Framework establishes streamlined and standardised program performance reporting to inform priority requirements. A small number of funded activities require additional mandatory data items to be reported. Go to Section 11 of this document for a comprehensive list of the field values.

5.3.1 Commonwealth Home Support Programme mandatory fields

The following items are required and will only present if the client is participating in the Commonwealth Home Support Programme activity:

- **Accommodation setting**: organisations are asked to record the accommodation setting category that best describes that of the client.
- **Living arrangements**: this is required for this program activity as it provides important information about a client’s presenting context. Living arrangements and its categories are adapted from the data collection definitions used by the AIHW. This information can also be collected as ‘household composition’ in the partnership approach.
- **DVA card status**: a client’s Department of Veterans’ Affairs (DVA) card status is collected.
- **Existence of a Caree**: this field is required to determine how many clients have care arrangements in place. This question is a yes/no response.
- **Amount of assistance provided**: measured as hours and minutes, quantity, cost and/or type. These data fields will only present once the service type is selected in the session. For more information go to the program specific guidance on the Data Exchange website and consult the CHSP Programme Manual.

- **Fees charged**: this item is captured at the session level. It allows organisations to report whether the participants of the session were charged a fee to attend the service and reflects the program activity policy regarding fee collection. This item is captured as a dollar figure.

- **Exit reason**: users can record the reason a client exited a service.

### 5.3.2 Family Law Services mandatory fields

The following items are required and will only present when a relevant Family Law Service is selected at the case creation level:

- **Parenting agreement reached**: this item is captured at the case level. It supports the measurement of an important program specific outcome, reflecting if a parenting agreement was reached between parties. The response options are Full agreement, Partial agreement, or No agreement.

- **Date of agreement**: records the date the parenting agreement was agreed or reached.

- **Did a legal practitioner assist with formalising agreement**: This yes/no question is also related to the above questions, indicating if a legal practitioner is present and participates in parenting mediation session. It must be completed.

- **Section 60(I) certificate type**: this item is used to record a section 60(I) certificate.

- **Date of certificate issued**: this item is related to the Section 60(I) certificate data field and records the date the Section 60(I) certificate was issued.

- **Fees charged**: this item is captured at the session level. It allows organisations to report whether the participants of the session were charged a fee to attend the service and reflects the program activity policy regarding fee collection. This item is captured as a dollar figure.

### 5.3.3 Financial Wellbeing and Capability mandatory field

The following items are only required for the Commonwealth Financial Counselling and Financial Capability, and Money Support Hubs program activities where it is part of the case creation process. When the service type ‘Education and Skills Training’ is selected, the following field becomes mandatory:

- **Money management course delivered**: this item is captured at the session level. The field contains the workshop types able to be delivered under this program activity and allows organisations to show that clients have undertaken the required training.

### 5.3.4 National Disability Advocacy Program mandatory fields

The following items are only required for the National Disability Advocacy Program (NDAP) program activities where it is part of the case creation process. The following question is mandatory:

- **Topic**: this item is captured at the session level. This field allows organisations to select the reason why a client is accessing a service.
6 Collecting partnership approach data

Organisations participating in the partnership approach report an extended data set, in exchange for access to additional self-service reports. Organisations can opt into the partnership approach, unless this is a requirement of their funding agreement.

Organisations who participate in the partnership approach are expected to report client circumstances SCOREs for the majority of their clients (over 50 per cent).

The extended data set includes information about a client’s presenting needs and circumstances, such as the reason for seeking assistance, referrals (in and out), household composition and income status. Other outcomes focused data is collected using Standard Client/Community Outcomes Reporting (SCORE).

In short to medium term service delivery, a SCORE is recorded in two parts: an initial SCORE (also known as a pre-SCORE) should be recorded against a session toward the beginning of service delivery; and a follow-up SCORE (also known as a post-SCORE) should be recorded against a session toward the end of that service delivery. In long term service delivery, multiple follow up SCOREs is recorded at regular intervals to track how the client’s outcomes change over time.

This section presents practical information about the extended data requirements of the partnership approach describing a client’s presenting needs and circumstances. Go to Section 7 of this document for detailed information about collecting and reporting SCORE and Section 11 for a list of data values.

6.1 Client needs and presenting context

Organisations participating in the partnership approach report additional data items about client needs and presenting circumstances where they already collect such data, or where they consider this information to be relevant and are able to collect it.

There are eleven data categories, set out below, which identify client needs and presenting circumstances. For many organisations, this information is already collected as part of internal service planning. Other items (such as migration visa) may only be relevant for specific funded activities. An organisation can choose to record some or all of the additional items, selecting those that are relevant to their clients and services.

All of the following listed items are optional.

6.2 Reasons for seeking assistance

Data about the reason clients sought assistance is collected to inform service planning to better respond to presenting needs. The optional categories for describing the reason for seeking assistance are standardised to reflect the SCORE outcome domains that cover the range of funded activities captured as part of the Data Exchange Framework. For each client, data is recorded about the main reason for seeking assistance and, if relevant, a secondary reason for seeking assistance. Reasons for seeking assistance are recorded at the case level, to allow organisations to reflect that clients go to different activities to address different needs. This field is optional.

The categories describing the reason for seeking assistance are:

- **Age-appropriate development**: where the client is seeking to improve age-appropriate development.
- **Community participation and networks**: where the client is seeking to change the impact of poor community participation and networks on their independence, participation and wellbeing.
- **Family functioning**: where the client is seeking to improve family functioning and change its impact on their independence, participation and wellbeing.
- **Financial Resilience**: where the client is seeking to improve financial resilience and change its impact to improve the client's independence, participation and wellbeing.
• **Employment**: to change the impact of a client’s lack of employment on their independence, participation and wellbeing.

• **Education and skills training**: where the client is seeking to engage with education and skills training to improve their independence, participation and wellbeing.

• **Material wellbeing and basic necessities**: to address the client’s immediate lack of money and basic items needed for day-to-day living and to improve their independence, participation and wellbeing.

• **Housing**: where the client is seeking to improve their housing stability or address the impact of poor housing on their independence, participation and wellbeing.

• **Mental health, wellbeing and self-care**: where the client is seeking to change the impact of mental health issues and self-care issues on their independence, participation and wellbeing.

• **Personal and family safety**: where the client is seeking to change the impact of personal and family safety issues on their independence, participation and wellbeing.

• **Physical health**: where the client is seeking to change the impact of their physical health on their independence, participation and wellbeing.

Many clients’ needs are complex and change over time. The ‘real’ reason for seeking assistance is often not apparent until after a client initially engages with the organisation. While recognising these limitations, data about the reasons for seeking assistance is recorded towards the start of the service to provide a high-level indication of the presenting need—within one of the standard circumstance outcome domains.

Reasons for seeking assistance is recorded as either the:

• **Primary reason for seeking assistance**: the main reason for seeking assistance, classified as one of ten possible categories.

• **Secondary reasons for seeking assistance**: the secondary reasons for seeking assistance, if relevant select another reason from the ten possible categories. In most cases this should be limited to one or two.

### 6.3 Referral source

Referral source is the person or agency responsible for referring a client to an organisation. The source of referral is important in mapping client pathways and access points. This optional field helps organisations to identify the main avenues their clients come through to reach their services. This information is used to help target networking and communication strategies to increase client engagement with a particular funded activity if desired. This field is optional.

Based on the data collection definitions used by the AIHW, referral source is classified into three standard categories (agency/organisation, non-agency, not stated). This information is recorded at the case level and allows organisations to reflect that clients are referred from different sources for each case.
6.4 Referrals to other services

Data is recorded about referrals made to other services to reflect a client’s need for help outside the scope of the funded activity. Referrals are recorded at the session level and can include one internal and one external type of referral per session record. Where more than one referral of each type is made, they are recorded in a separate session.

Two data items are recorded to reflect referrals to other services:

Referral to other service
- **Internal**: made to another service offered within the same organisation
- **External**: made to a service that is provided by a different organisation

Referral purpose
- **Physical health**: the client is referred to assist with the impact of their physical health on their independence, participation and wellbeing.
- **Mental health wellbeing and self-care**: the client is referred to help the impact of client’s mental health and self-care issues on their independence, participation and wellbeing.
- **Personal and family safety**: the client is referred to help with the impact of personal and family safety issues on their independence, participation and wellbeing.
- **Age-appropriate development**: the client is referred to help improve age-appropriate development.
- **Community participation and networks**: the client is referred to help with the impact of poor community participation and networks on their independence, participation and wellbeing.
- **Family functioning**: the client is referred to improve family functioning and change its impact to improve the client’s independence, participation and wellbeing.
- **Financial Resilience**: the client is referred to help improve financial resilience and change its impact to improve the client’s independence, participation and wellbeing.
- **Employment**: the client is referred to help with the impact of a client’s lack of employment on their independence, participation and wellbeing.
- **Education and skills training**: the client is referred to help with the impact of a client’s inability to engage with education and skills training on their independence, participation and wellbeing.
- **Material wellbeing and basic necessities**: the client is referred to help with the impact of the client’s immediate lack of money and basic items needed for day-to-day living to improve their independence, participation and wellbeing.

- **Housing**: the client is referred to improve their housing stability or address the impact of poor housing on their independence, participation and wellbeing.

- **Support to caring role**: the client is referred to help with their caring responsibilities.

- **Other**: the referral purpose is not captured in the list provided.

### 6.5 Household composition

Data is reported about household composition as it provides important information about a client’s presenting context. Household composition is classified into standard categories that are adapted from the data collection definitions used by AIHW. Household composition is adapted to include data on whether the client is homeless. This field is optional.

### 6.6 Highest level of education/qualification

Data is reported about highest educational achievement a person has attained. It lists qualifications and other educational attainments regardless of the particular field of study or the type of institution in which the study was undertaken. This field is optional.

### 6.7 Employment status

Data is reported about a client’s employment status. Employment status is classified into standard categories that are adapted from the data collection definitions used by Disability Employment Services. It is important to note a client’s employment status can also include being a carer.

### 6.8 Household income

#### 6.8.1 Main source of income

Data is reported about main source of income to provide important information about a client’s presenting needs. Main source of income is classified into standard categories, based on the data collection definitions used by the AIHW. This field is optional. Go to Section 11 for a list of these categories.

#### 6.8.2 Approximate gross income

Data is reported about approximate income as it provides important information about a client’s presenting context. Data is recorded as a numerical value for the estimated dollar amount of the client’s gross income (earnings before tax). This is based on what is reported by the client and no evidence of income is required.

As clients can receive their income at different frequencies, the Data Exchange web-based portal allows estimated gross income to be provided as either a weekly, fortnightly, monthly or annual figure. Income frequency must be recorded in order to be able to record the approximate gross income. This field is optional.

### 6.9 Expanded CALD indicators

Optional data is reported about CALD background as this information provides important context about clients’ circumstances. This includes:

- **Date of first arrival in Australia**: records the date a client first arrived in Australia where appropriate for a funded activity. This field is primarily included as an eligibility indicator for the Settlement Engagement and Transition Support (SETS) Program. It can also be used as a potential indicator of disadvantage for other program activities.

- **Migration Visa category**: records the Visa category the client arrived on, where appropriate for a funded activity. This field is primarily included as an eligibility indicator for the SETS Program. It can also be used as a potential indicator of disadvantage for other program activities.
- **Ancestry**: records a client’s ancestry if relevant. The list of values is drawn from the Australian Bureau of Statistics [Australian Standard Classification of Cultural and Ethnic Groups (ASCCEG), 2016](https://www.abs.gov.au).  

6.10 **Homeless indicator**  
Data is reported about a client’s housing situation. Noting the values for the homeless indicator are Yes, No or At Risk, a person is homeless if they do not have suitable accommodation alternatives and their current living arrangement:

- is in a dwelling that is inadequate;
- has no tenure, or if their initial tenure is short and not extendable; or
- does not allow them to have control of, and access to space for social relations

A person may be at risk of homelessness in a number of situations including living in housing with major structural problems, residents are in constant threat of violence, living in crowded or improvised dwellings, or persons who are marginally housed in caravan parks.

The response should be based solely on what is reported by the client. This field is optional.

6.11 **Attendance profile**  
Data is reported to better understand the relationship between clients within a case. This should be based on the relationships self-reported by the client and other attendees. The different data items that can be selected are family, community event, peer support group, couple and cohabitants.

In the Data Exchange a family is defined as two or more people, one of whom is at least 15 years of age, who are related by blood, marriage (registered or de facto), adoption, step or fostering, and who are usually resident in the same household.

A couple is defined as two people usually residing in the same household who share a social, economic and emotional bond who consider their relationship to be a marriage or marriage like union. This relationship is identified by the presence of a registered marriage or de facto relationship.

Cohabitants are people in residence together who are not related or in domestic relationship. Peer support groups are a planned and structured service.

A community event is a large-scale, locality based event where attendees generally do not register and are not identified. This field is optional.

6.12 **Carer Status**  
Data is reported on the client’s self-report status as a carer. A carer is defined as a person who provides unpaid care and support to family members and friends who have a disability, mental illness, chronic condition, terminal illness, an alcohol or other drug issue or who are frail aged. Data in this field is based on the client’s reported caring situation. This field is optional.

6.13 **NDIS eligibility**  
Data is reported on the NDIS eligibility of a client, and if they have an application in progress. This field is optional.

6.14 **Service setting**  
Data is reported to help differentiate where services are provided. The service setting categories in the Data Exchange are:

- **Organisation outlet/office**: the organisation’s outlets as recorded in the Data Exchange
- **Client’s residence**: the client’s usual place of residence. Please refer to the section residential address for more information
- **Community venue**: a venue that is available to the general public and is hired away from the organisation’s usual offices. Examples include community halls, public libraries or parks

- **Partner organisation**: another organisations facilities (whether there is a formal or informal arrangement) to deliver a service

- **Healthcare facility**: doctor’s office, hospital, mental health facility, aged care facility

- **Education facility**: school, university

- **Justice Facility**: correctional facility (jail, prison), police station, court of law

- **Telephone**: person to person contact (other than face to face) that are interactive. Examples includes services provided to clients over the phone, via video conference or online chat

- **Digital**: self-service, e-Learning, webinar, online training, non-interactive electronic resources.

### 6.15 Interpreter present

Data is reported on whether an interpreter was present for the instance of service to assist with translation and facilitation. This could be an interpreter provided by the organisation or someone the client has brought along to help them. This field is optional.
7 Recording client and community SCOREs

The main focus of the extended data set (the partnership approach) is collecting information about client and community outcomes achieved for individuals accessing funded activities. Client and community outcomes are achieved in different ways and progressively over different periods of time. They range from immediate short term outcomes to longer-term changes that positively affect a client’s life circumstances.

SCORE is designed to captures outcomes consistently, in ways that do not impose additional administrative costs on organisations, recognising that organisations are not funded to be specialist researchers or to spend disproportionate amounts of time measuring outcomes.

Many organisations use a diverse range of instruments and methods for measuring and assessing client and community outcomes, often linked to organisational and sector specific priorities. The SCORE tool used in the partnership approach allows organisations to measure outcomes using existing validated tools, but report them through SCORE so as to be captured in a consistent and comparable way.

In some program areas, specific validated instruments are already used (such as Kessler 10). A translation matrix was developed to help organisations convert results from commonly used outcomes measurement tools into SCORE.

If the organisation does not have a standard or systematic approach to measuring outcomes, they can use SCORE as an outcomes measurement tool. Alternatively, organisations may prefer to adapt the SCORE scales and domains to create their own generic outcomes measurement tool that is adjusted for their service.

There are four different types of outcomes measured through SCORE that help tell the story of what was achieved; three for individual clients (their circumstances, goals and satisfaction) and one for a group/community:

- **Circumstances**: changes in client circumstances, such as mental/physical health, material wellbeing and situation.
- **Goals**: progress in achieving specific goals, such as behaviours, skills for lessening the impact of a crisis.
- **Satisfaction**: did the client feel the service met their needs?
- **Community**: changes in group, organisation, and community capacity to address identified needs.

SCOREs are captured at the case level, and are reported using a five point rating scale. This provides a consistent and comparable way to translate outcomes across programs using the Data Exchange. A SCORE is also recorded in two parts; using an initial SCORE towards the beginning of service and a subsequent SCORE either at the end of service delivery, or at regular intervals into the future to track a client’s progress. Go to the Data Exchange website for program specific guidance information on SCORE and the Translation Matrix.

7.1 How to record into SCORE

Organisations have the flexibility to record outcomes directly into SCORE, or to use externally validated instruments or internally developed tools. When reporting this information into the Data Exchange, the ‘Assessed by’ field captures whether a validated tool was used, and who made the assessment.

A validated instrument is psychometrically tested for reliability (the ability of the instrument to produce consistent results), validity (the ability of the instrument to produce true results), and sensitivity (the probability of correctly identifying a client with the condition).

If the organisation doesn’t have an outcomes measurement tool, or have developed their own tool but have not validated it, the outcomes data should be reported as using ‘SCORE directly’ in the ‘Assessed by’ field.

A SCORE may be determined by the practitioner’s professional assessment, a client’s self-assessment, a joint assessment between the client and practitioner, or an assessment by the client’s support person (such
as a carer). All of these assessments are conducted using the SCORE tool, or outcomes tools validated instruments. While they are voluntary fields, it is recommended organisations report who conducted the assessment and the type of tool used.

Organisations are encouraged to collect SCORE in a way that bests suits their own unique service delivery context.

7.2 Collecting and reporting Circumstances SCOREs

Organisations who participate in the partnership approach agree to report an initial and at least one subsequent Circumstances SCORE of the relevant domain(s) for the majority of their clients (over 50 per cent).

It is valuable to have multiple SCOREs for all clients however, it is recognised that there are a range of situations where this may not be possible. For example, subsequent SCOREs may not be able to be recorded due to clients unexpectedly exiting a service.

7.2.1 Circumstance domains

The Circumstance SCORE is linked to eleven outcome domains. The type of outcome domain that is relevant to each client depends on the context of the funded activity being delivered. Organisations should only report a numerical rating against the domain(s) that are most relevant for their client and the outcomes they are aiming to achieve.

There are potential connections between all of the domains across program activities and there are no fixed rules for selecting the most relevant domain for a particular client or funded activity. However, guidance on which domains are most relevant is provided by policy areas of each program using the Data Exchange.

The Circumstance domain descriptions are:

- **Physical health**: the funded activity is seeking to change the impact of a client’s physical health to improve their independence, participation and wellbeing.

- **Mental health, wellbeing and self-care**: the funded activity is seeking to change the impact of a client’s mental health and self-care, to improve their independence, participation and wellbeing.

- **Personal and family safety**: the funded activity is seeking to change the impact of personal and family safety issues to improve the client’s independence, participation and wellbeing.

- **Age-appropriate development**: the funded activity is seeking to improve a client’s age-appropriate development to improve the client’s independence, participation and wellbeing.

- **Community participation and networks**: the funded activity is seeking to change the impact of poor community participation and networks to improve a client’s independence, participation and wellbeing.

- **Family functioning**: the funded activity is seeking to improve family functioning and change its impact so it enhances the family’s independence, participation and wellbeing.

- **Financial Resilience**: the funded activity is seeking to change the impact of poor money management to improve the client’s independence, participation and wellbeing.

- **Employment**: the funded activity is seeking to change the impact of a client’s lack of employment on their independence, participation and wellbeing.

- **Education and skills training**: the funded activity is seeking to change the impact of a client’s ability to engage with education and skills training on their independence, participation and wellbeing.

- **Material wellbeing and basic necessities**: the funded activity is seeking to change the impact of the client’s immediate lack of money and basic items needed for day-to-day living to improve their independence, participation and wellbeing.
- **Housing:** the funded activity is seeking to improve the client’s housing stability or address the impact of poor housing to improve their independence, participation and wellbeing.

### 7.2.2 Circumstance rating scale

To record a client SCORE, organisations need to record a rating between 1 and 5 against a relevant domain. A five point rating scale is used to report changes in client outcomes when using SCORE. This scale is used for all four SCORE components (Circumstance, Goal, Satisfaction and Community):

1. is used to report that the client’s current circumstances are having a significant *negative impact* on their independence, participation and wellbeing. For example, a significant negative impact may be a lack of safe housing on an individual’s independence, or poor family functioning that impacts on the client’s wellbeing.

2. is used to report that the client’s current circumstances are having a *moderate negative impact* on their independence, participation and wellbeing—for example, a moderate negative impact of poor physical health that impacts on their independence; or a moderate negative impact of family functioning on the client’s wellbeing.

3. is used to report that the client’s current circumstances are in a ‘middle ground’ between the significant/moderate negative impacts (indicated by Ratings 1 and 2) and the adequate circumstances over the short and medium term (indicated by Ratings 4 and 5). For example, progress towards improving a client’s community participation without having reached a point where the client’s current circumstances are adequate over the short or medium term.

4. is used to report that the client’s current circumstances are *adequate over the short term* to support their independence, participation and wellbeing—for example, adequate access to money to meet basic needs in the short-term; adequate family functioning to support the family’s wellbeing in the short-term.

5. is used to report that the client’s current circumstances are *adequate and stable over the medium term* to support their independence, participation and wellbeing—for example, adequate and stable money management to support an individual’s independence; adequate and stable family functioning to support the family’s wellbeing.

A summary of the ‘generic’ scale for the Circumstance SCORE is:
## Table 3. Circumstances SCORE domains

<table>
<thead>
<tr>
<th>circumstance SCORE domain</th>
<th>1: Negative Impact</th>
<th>2: Moderate negative impact</th>
<th>3: Middle ground</th>
<th>4: Adequate over the short term</th>
<th>5: Adequate and stable over the medium term</th>
</tr>
</thead>
<tbody>
<tr>
<td>Physical health</td>
<td>Significant negative impact of poor physical health on independence, participation and wellbeing</td>
<td>Moderate negative impact of poor physical health on independence, participation and wellbeing</td>
<td>Progress towards improving physical health to support independence, participation and wellbeing</td>
<td>Sustained initial improvements in physical health to support independence, participation and wellbeing</td>
<td>Adequate ongoing physical health to support independence, participation and wellbeing</td>
</tr>
<tr>
<td>Mental health, wellbeing and self-care</td>
<td>Significant negative impact of poor mental health, wellbeing and self-care on independence, participation and wellbeing</td>
<td>Moderate negative impact of poor mental health, wellbeing and self-care on independence, participation and wellbeing</td>
<td>Progress towards improving mental health, wellbeing and self-care to support independence, participation and wellbeing</td>
<td>Adequate short-term mental health, wellbeing and self-care to support independence, participation and wellbeing</td>
<td>Adequate ongoing mental health, wellbeing and self-care to support independence, participation and wellbeing</td>
</tr>
<tr>
<td>Personal and family safety</td>
<td>Significant negative impact of poor personal and family safety on independence, participation and wellbeing</td>
<td>Moderate negative impact of poor personal and family safety on independence, participation and wellbeing</td>
<td>Progress towards improving personal and family safety to support independence, participation and wellbeing</td>
<td>Adequate short-term personal and family safety to support independence, participation and wellbeing</td>
<td>Adequate ongoing personal and family safety to support independence, participation and wellbeing</td>
</tr>
<tr>
<td>Age-appropriate development</td>
<td>Significant negative impact of poor age-appropriate development on independence, participation and wellbeing</td>
<td>Moderate negative impact of poor age-appropriate development on independence, participation and wellbeing</td>
<td>Progress towards improving age-appropriate development to support independence, participation and wellbeing</td>
<td>Adequate short-term age-appropriate development to support independence, participation and wellbeing</td>
<td>Adequate ongoing age-appropriate development to support independence, participation and wellbeing</td>
</tr>
<tr>
<td>Community participation and networks</td>
<td>Significant negative impact of poor community participation and networks on independence, participation and wellbeing</td>
<td>Moderate negative impact of poor community participation and networks on independence, participation and wellbeing</td>
<td>Progress towards improving community participation and networks to support independence, participation and wellbeing</td>
<td>Adequate short-term community participation and networks to support independence, participation and wellbeing</td>
<td>Adequate ongoing community participation and networks to support independence, participation and wellbeing</td>
</tr>
<tr>
<td>Family functioning</td>
<td>Significant negative impact of poor family functioning on independence, participation and wellbeing</td>
<td>Moderate negative impact of poor family functioning on independence, participation and wellbeing</td>
<td>Progress towards improving family functioning to support independence, participation and wellbeing</td>
<td>Adequate short-term family functioning to support independence, participation and wellbeing</td>
<td>Adequate ongoing family functioning to support independence, participation and wellbeing</td>
</tr>
</tbody>
</table>
7.3 Collecting and reporting Goals SCOREs

Organisations who participate in the partnership approach agree to report an initial and at least one subsequent Goals SCORE of the relevant domain(s) for the majority of their clients (over 50 per cent).

It is valuable to have multiple SCOREs for all clients however, it is recognised that there are a range of situations where this may not be possible. There are occasions where follow-up SCORES may not be able to be recorded due to clients unexpectedly exiting a service.

A client Goals SCORE is a measure of the extent of achievement of a client’s individual goals. The term ‘goal’ is used very broadly. In some service contexts, goals refer to planned objectives that are documented...
in the client’s case plan. In other situations, goals are better described as simply the things clients want help with. In both scenarios, the client Goals SCORE is used to measure the goal attainment or extent to which the organisation is able to address the things a client sought help with.

7.3.1 Goals domains

The Goals SCORE is linked to six outcome domains. The type of outcome domain relevant to each client depends on the context of the goals a client wants to achieve. Organisations should only report a rating against the domain(s) that are most relevant for their client and the outcomes they are aiming to achieve.

As there are potential connections between all of the domains and program activities, there are no fixed rules for selecting the most relevant domain for a particular client or funded activity. However, guidance on the most relevant domains is provided by policy areas of each program using the Data Exchange.

The goal domain descriptions are:

- **Changed knowledge and access to information**: the funded activity is seeking to change a client’s knowledge and understanding of issues to improve their independence, participation and wellbeing or to improve their access to relevant information about these issues.
- **Changed skills**: the funded activity is seeking to enhance a client’s skills set to improve their independence, participation and wellbeing.
- **Changed behaviours**: the funded activity is seeking to change a client’s behaviours to improve their independence, participation and wellbeing.
- **Empowerment, choice and control to make own decisions**: the funded activity is seeking to enhance a client's confidence, choice and control to make their own decisions and take empowerment on issues that impact on their independence, participation and wellbeing.
- **Engagement with relevant support services**: the funded activity is seeking to improve a client’s engagement with services to support their independence, participation and wellbeing.
- **Changed impact of immediate crisis**: the funded activity is seeking to address or reduce the impact of an immediate crisis to improve the client’s independence, participation and wellbeing.

7.3.2 Goals rating scale

A five point rating scale is used to report changes in client outcomes when using SCORE. This scale is used for all four SCORE components (Circumstances, Goals, Satisfaction and Community):

1: is used to report that the client has made **no progress** in achieving their individual goals within the selected goal domain.
2: is used to report that the client has made **limited progress** in achieving their individual goals within the selected goal domain but there is emerging engagement in addressing the issues they sought assistance with.
3: is used to report that the client has made **limited progress** in achieving their individual goals within the selected goal domain but there is **strong engagement** in addressing the issues they sought assistance with.
4: is used to report that the client has made **moderate progress** in achieving their individual goals within the selected goal domain.
5: is used to report that the client has **fully achieved** their individual goals within the selected outcomes domain.

To record a client SCORE, organisations need to record a rating between 1 and 5 against a relevant domain. A summary of the ‘generic’ scale for the Goals SCORE is:
Table 4. Goals SCORE domains

<table>
<thead>
<tr>
<th>Goal SCORE domain</th>
<th>1: No progress</th>
<th>2: Limited progress with emerging engagement</th>
<th>3: Limited progress with strong engagement</th>
<th>4: Moderate progress</th>
<th>5: Fully achieved</th>
</tr>
</thead>
<tbody>
<tr>
<td>Knowledge and access to information</td>
<td>No progress in increasing access to information and knowledge in areas relevant to clients' needs and circumstances</td>
<td>Limited progress to date in achieving information/ knowledge goals–but emerging engagement</td>
<td>Limited progress to date in achieving information/ knowledge goals–but strong engagement</td>
<td>Moderate progress to date in achieving information/ knowledge goals</td>
<td>Full achievement of goals related to increasing access to information and knowledge in areas relevant to client's needs and circumstances</td>
</tr>
<tr>
<td>Skills</td>
<td>No progress in increasing skills in areas relevant to client’s needs and circumstances</td>
<td>Limited progress to date in achieving skills goals–but emerging engagement</td>
<td>Limited progress to date in achieving skills goals–but strong engagement</td>
<td>Moderate progress to date in achieving skills goals</td>
<td>Full achievement of goals related to increasing skills in areas relevant to client's needs and circumstances</td>
</tr>
<tr>
<td>Behaviours</td>
<td>No progress in changing behaviours in areas relevant to client’s needs and circumstances</td>
<td>Limited progress to date in achieving behaviour goals–but emerging engagement</td>
<td>Limited progress to date in achieving behaviour goals–but strong engagement</td>
<td>Moderate progress to date in achieving behaviour goals</td>
<td>Full achievement of goals related to changing behaviours in areas relevant to client's needs and circumstances</td>
</tr>
<tr>
<td>Empowerment, choice and control to make own decisions</td>
<td>No progress in increasing confidence and exercising choice/control in making decisions that impact client’s needs</td>
<td>Limited progress to date in achieving empowerment, choice and control goals–but emerging engagement</td>
<td>Limited progress to date in achieving empowerment, choice and control goals–but strong engagement</td>
<td>Moderate progress to date in achieving empowerment, choice and control goals</td>
<td>Full achievement of goals related to increasing confidence and exercising choice/control in making decisions that impact client’s needs</td>
</tr>
<tr>
<td>Engagement with support services</td>
<td>No progress in increasing engagement with support services relevant to client’s needs and circumstances</td>
<td>Limited progress to date in achieving engagement goals – but emerging engagement</td>
<td>Limited progress to date in achieving engagement goals–but strong engagement</td>
<td>Moderate progress to date in achieving engagement goals</td>
<td>Full achievement of goals related to increasing engagement with support services relevant to client’s needs and circumstances</td>
</tr>
<tr>
<td>Impact of immediate crisis</td>
<td>No progress in reducing the negative impact of the immediate crisis</td>
<td>Limited progress to date in achieving goals to reduce the negative impact–but emerging engagement</td>
<td>Limited progress to date in achieving goals to reduce the negative impact–but strong engagement</td>
<td>Moderate progress to date in achieving goals to reduce the negative impact</td>
<td>Full achievement of goals related to reducing the negative impact of the immediate crisis</td>
</tr>
</tbody>
</table>
7.4 Collecting and reporting client Satisfaction SCOREs

Unlike the other areas of SCORE, satisfaction is only measured at the end of service delivery. Organisations who participate in the partnership approach agree to report client Satisfaction SCOREs for a small sample of their clients (at least 10 per cent per reporting period).

It is valuable to have Satisfaction SCOREs for all clients, however this may be time consuming for some organisations to collect. There can also be situations where SCOREs cannot be collected, such as clients unexpectedly exiting a service.

7.4.1 Satisfaction domains

The Satisfaction SCORE relates to three key questions about a client’s perceptions of the responsiveness and value of the service received:

- The service listened to me and understood my issues
- I am satisfied with the services I have received
- I am better able to deal with issues that I sought help with.

7.4.2 Satisfaction rating scale

A five point rating scale is used to report changes in client outcomes when using SCORE. This scale is used for all four SCORE components (Circumstances, Goals, Satisfaction and Community):

1: is used to report that the client **disagrees** with the statement about the service (for example, disagrees that the service listened to them and understood their issues).

2: is used to report that the client **tends to disagree** with the statement about the service (for example, tends to disagree that the service listened to them and understood their issues).

3: is used to report that the client **neither agrees or disagrees** with the statement about the service (for example, neither agrees nor disagrees that the service listened to them and understood their issues).

4: is used to report that the client **tends to agree** with the statement about the service (for example, tends to agree that the service listened to them and understood their issues).

5: is used to report that the client **agrees** with the statement about the service (for example, agrees that the service listened to them and understood their issues).

To record a client SCORE, organisations need to record a rating between 1 and 5 against a relevant domain.
Table 5. Satisfaction SCORE domains

<table>
<thead>
<tr>
<th>Satisfaction SCORE domain</th>
<th>1: Disagree</th>
<th>2: Tend to disagree</th>
<th>3: Neither agree or disagree</th>
<th>4: Tend to agree</th>
<th>5: Agree</th>
</tr>
</thead>
<tbody>
<tr>
<td>The service listened to me and understood my issues</td>
<td>Disagrees that the service listened to me and understood my issues</td>
<td>Tend to disagree that the service listened to me and understood my issues</td>
<td>Neither agrees nor disagrees that the service listened to me and understood my issues</td>
<td>Tends to agree that the service listened to me and understood my issues</td>
<td>Agrees that the service listened to me and understood my issues</td>
</tr>
<tr>
<td>I am satisfied with the services I have received</td>
<td>I am not satisfied with the services I have received</td>
<td>Tends to disagree that I was satisfied with the services I have received</td>
<td>Neither agrees nor disagrees that the services listened to me and understood my issues</td>
<td>Tends to agree that I was satisfied with the services I have received</td>
<td>I am satisfied with the services I have received</td>
</tr>
<tr>
<td>I am better able to deal with issues that I sought help with</td>
<td>Disagrees that I am better able to deal with my issues</td>
<td>Tends to disagree that I am better able to deal with my issues</td>
<td>Neither agrees nor disagrees that I am better able to deal with my issues</td>
<td>Tends to agree that I am better able to deal with my issues</td>
<td>Agrees that I am better able to deal with my issues</td>
</tr>
</tbody>
</table>

7.4.3 Measurement and reporting of client Satisfaction SCOREs

If an organisation already uses an existing outcomes measurement tool that meets their needs, they can continue to use it and translate the outcome data to SCORE.

If an organisation does not currently have a standard or systematic approach to measure client satisfaction, it is possible to use the Satisfaction SCORE table above as a simple survey tool. Alternatively, organisations can adapt the SCORE scales and domains to create their own interim measurement tool that is adjusted to the organisation’s service type.

Given that Satisfaction SCOREs are intended to be collected as part of good practice service delivery, each organisation is responsible for ensuring that the data collection process is ethical and reliable. Key practices to consider when collecting client feedback are outlined below.

7.4.4 Data collection methods

It is important that client satisfaction feedback is voluntary and that it does not cause discomfort or anxiety for clients. In particular, the pre-existing relationship between clients and the organisation may bias client results. A number of measures are taken to ensure that clients understand the nature of participating and their options:

- All participants are provided with information that makes it clear that completing the survey is voluntary.
- Participants are provided with information that makes it clear that the purpose of collecting the client feedback is to improve services.
- Participants should be encouraged to share their honest views and to make constructive suggestions if they think the service could be improved.

7.4.5 Client confidentiality

It is important that clients can provide feedback anonymously. There is a range of ways to achieve this, for example:

- The client feedback is collected by a person not directly involved in service delivery.
- Using a survey so that clients can complete and submit without showing the organisation their response.
- If paper-based forms are used, consider using sealed envelopes for returning completed surveys.

7.5 Collecting and reporting Community SCOREs

Organisations who participate in the partnership approach agree to report Community SCOREs for the majority of group or community activities where it is not feasible to record the changes for individual clients.

7.5.1 Community domains

The Community SCORE is linked to four domains that reflect changes that may occur for a group or community rather than individual clients:

- Group/Community knowledge, skills, attitudes and behaviours for a group of clients or community members participating in the service (where it is not feasible to record the changes for individual members of the group or community)
- Organisational knowledge, skills and practices to better respond to the needs of targeted clients or communities.
- Community infrastructure and networks to better respond to the needs of targeted clients and communities.
- Social cohesion to demonstrate greater community cohesion and social harmony.

7.5.2 Community rating scale

A five point rating scale is used to report changes in community outcomes when using SCORE. This scale is used for all four SCORE components (Circumstances, Goals, Satisfaction and Community).

The community domain descriptions are:

1: is used to report **no change** in the group/organisation/community capacity to address or respond to their own needs or the needs of targeted clients and communities.
2: is used to report **limited change** in the group/organisation/community capacity to address or respond to their own needs or the needs of targeted clients and communities, but there is **emerging engagement** in addressing the issues.
3: is used to report **limited change** in the group/organisation/community capacity to address or respond to their own needs or the needs of targeted clients and communities, but there is **strong engagement** in addressing the issues.
4: is used to report **moderate change** in the group/organisation/community capacity to address or respond to their own needs or the needs of targeted clients and communities.
5: is used to report **significant change** in the group/organisation/community capacity to address or respond to their own needs or the needs of targeted clients and communities.

To record a Community SCORE, organisations need to record a rating between 1 and 5 against a relevant domain. A summary of the ‘generic’ scale for the Community SCORE is:
Table 6. Community SCORE domains

<table>
<thead>
<tr>
<th>Community SCORE domain</th>
<th>1: No change</th>
<th>2: Limited change with emerging engagement</th>
<th>3: Limited change with strong engagement</th>
<th>4: Moderate change</th>
<th>5: Significant change</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group/community knowledge, skills, attitudes behaviours</td>
<td>No change in knowledge, skills, attitudes, behaviours</td>
<td>Limited change in knowledge, skills, attitudes, behaviours–but emerging engagement</td>
<td>Limited change in knowledge, skills, attitudes, behaviours–but strong engagement</td>
<td>Moderate change in knowledge, skills, attitudes, behaviours</td>
<td>Significant positive change in knowledge, skills, attitudes, behaviours</td>
</tr>
<tr>
<td>Organisational knowledge, skills and practices</td>
<td>No change in organisational knowledge, skills, practices to respond to the needs of targeted clients/communities</td>
<td>Limited change in organisational knowledge, skills, practices–but emerging engagement</td>
<td>Limited change in organisational knowledge, skills, practices–but strong engagement</td>
<td>Moderate change in organisational knowledge, skills, practices</td>
<td>Significant positive change in organisational knowledge, skills, behaviours to better respond to the needs of targeted clients/communities</td>
</tr>
<tr>
<td>Community infrastructure and networks</td>
<td>No change in community infrastructure/networks to respond to the needs of targeted clients/communities</td>
<td>Limited change in community infrastructure/networks–but emerging engagement of community networks</td>
<td>Limited change in community infrastructure/networks–but strong engagement of community networks</td>
<td>Moderate change in community infrastructure/networks</td>
<td>Significant positive change in community infrastructure/networks to better respond to the needs of targeted clients/communities</td>
</tr>
<tr>
<td>Social cohesion</td>
<td>No change in demonstration of greater community cohesion and social harmony</td>
<td>Limited change in demonstration of greater community cohesion and social harmony–but emerging engagement in issues</td>
<td>Limited change in demonstration of greater community cohesion and social harmony–but stronger engagement in issues</td>
<td>Moderate demonstration of greater community cohesion and social harmony</td>
<td>Significant positive demonstration of greater community cohesion and social harmony</td>
</tr>
</tbody>
</table>

7.5.3 Measurement and reporting of Community SCORES

If an organisation already uses an existing outcomes measurement tool that meets their needs, they can continue to use it and simply translate the outcome data to SCORE. For example, if an organisation is using an outcomes measurement tool that is used and reported on by other evaluators and researchers to measure changes in community circumstances on a numerical scale, they can create a ‘concordance’ table to report these outcomes as part of the Data Exchange. Go to the Data Exchange website for more information on the Translation Matrix.
8 Program specific surveys

As the host of the Data Exchange, the department is looking at additional ways to better understand how funded services are achieving outcomes for individuals and communities.

Asking clients if they would like to participate in follow-up research, such as surveys and evaluation, forms part of the Data Exchange priority requirements—the standardised core set of mandatory reporting that applies to all Data Exchange in-scope programs.

Following a survey pilot in 2016 and extensive consultations with organisations and other stakeholders, the department has decided to implement program specific surveys, rather than a generalised outcomes survey for all Data Exchange clients.

8.1 Current surveys

In 2018, the department commenced a program specific survey for the Try Test and Learn Fund. All organisations funded to deliver the Try Test and Learn Fund are required to offer clients the program specific survey, as part of the Try Test and Learn Fund evaluation.

Go to the Data Exchange website for the latest information about program specific surveys and follow-up research.
9 Data Exchange reports

As part of the Data Exchange, all organisations that use the Data Exchange will have access to their own set of reports, which reflect the information submitted by their organisation. All available reports are accessed via the Data Exchange web-based portal. The ability to access the data and run reports will reflect the level of user access within the organisation.

Go to the Data Exchange website for detailed information on this topic and access to related training resources.

9.1 Report types

Standard self-service reports
These reports cover the mandatory priority data submitted by the organisation during a particular reporting period. For a current open reporting period the report will refresh every 24 hours to allow near real-time access to the information transmitted.

Partnership approach reports
Organisations participating in the partnership approach have access to a sophisticated suite of additional reports. Using both priority requirement data and extended partnership data, combined with government and population data sets, these reports provide valuable insights into service delivery and client outcomes.

Client survey reports
Reports covering aggregate client survey data will be developed for program specific surveys as needed.

9.2 Benefits of reports

Reports make the data is entered visible and enables verification of data quality and integrity. They also provide organisations with an evidence base for evaluation and to inform best practice. The Data Exchange uses de-identified, aggregate information to look at both short and long term outcomes achieved for clients across the broad suite of in-scope programs. The reports allow for an understanding of the collective impact of service provided and what combinations of services deliver the best outcomes for clients.

9.3 Access and visibility of reports

Within the Data Exchange, access and visibility of reports will depend on the way organisations set up their outlets and delivery partners.

By default, organisations cannot see a delivery partner's data. However, the ‘handshake’ allows the sharing of reports data in the form of de-identified, aggregate information. The handshake is a virtual agreement between a lead organisation and their delivery partner(s), to share data from the delivery partner to the lead organisation for their activity. Under a handshake, a lead organisation can only access data reported by the delivery partner for the agreed program(s).
10 Administrative matters

10.1 In-scope program activities for the Data Exchange

A list of program activities in-scope for the Data Exchange is on the Data Exchange website. This list is updated on a regular basis as new program activities start using the Data Exchange.

10.2 Access and set-up

In order to use the Data Exchange, an organisation must complete a number of access and set-up steps before client and session information is entered into the system. Organisations are strongly encouraged to complete these steps as early as possible in the reporting period. They include:

- applying for an AUSkey through the Australian Business Register
- submitting a User Access Request to the Data Exchange Helpdesk
- accessing the Data Exchange portal to set up their organisation
- create Outlets
- add program activities to Outlets
- add delivery partner details (if required)
- create additional users (if required)
- setting up bulk upload (if required).

Go to the ‘Quick Start Guide’ and other training and guidance materials on the Data Exchange website.

Completing access and set-up steps in a timely manner is the responsibility of the organisation as part of their funding agreement obligations.

If these steps are completed too close to the end of a reporting period, the department may not be able to process access and set-up requests with sufficient time remaining for the organisation to complete their data reporting before the due date.

10.3 Reporting periods and deadlines

The Data Exchange has two standardised six monthly performance reporting periods each year:

Users of the Data Exchange have an extra 30 days at the end of each reporting period, known as the ‘close-off period’, to allow time to quality check their data and make amendments to reported data. After the 30 day close-off period the Data Exchange automatically closes and no longer accepts uploads for that reporting period.
Organisations can enter data at any time within a reporting period, and are encouraged to do so regularly to make best use of the self-service reports and avoid unnecessary backlog or ‘crunch’ periods. Organisations new to the Data Exchange, in particular, need to plan for and allow sufficient time for access, set-up and other lead times, in order to meet reporting deadlines.

Once a reporting period has closed, data relating to that period of time will no longer be able to be recorded. Data outside of a reporting period may only be entered if an organisation has sought and is granted a system re-opening.

10.4 Compliance issues and system re-open requests

If an organisation experiences a crisis or event outside of their control that will impact their ability to meet performance reporting requirements, they can request a re-opening of the system.

System re-opening requests are submitted via the ‘Request to re-open the Data Exchange form’ on the Data Exchange website, however organisations should also consult with their Funding Arrangement Manager or funding agency contact.

System re-openings will only be granted under exceptional circumstances following consultation with Funding Arrangement Managers. Submission of a request does not guarantee a system re-opening will be granted.

10.5 Flexible ways to transmit data

Users can transmit their data to the Data Exchange in one of three ways; system-to-system transfer, bulk file upload, or manual entry into the web-based portal. It is recommended to select one of these as the main transmission method for the longer term. However, in some circumstances, such as the period of initial transition into the Data Exchange, manual entry may need to be used in combination with another transmission method.

All users of the Data Exchange must have an AUSkey account before registering for the system. AUSkey is a secure credential and is used for many government online services on behalf of a business or entity. All government agencies are required to implement AUSkey for all new business to government transactions, and many organisations already have AUSkeys for other purposes. AUSkey is administered by the Australian Business Register (ABR). Go to the ABR website for more information.

At least one person within each organisation will need to complete and submit the Data Exchange User Access Request Form to have Org Administrator access to the Data Exchange. This form is on the Data Exchange website.

10.5.1 System-to-system transfers

Organisations with their own client management software systems capable of pushing data via web services through to the Data Exchange can continue using this software to collect and transfer their performance data. Organisations will need to make a one-off adjustment (or ‘enhancement’) to their application in accordance with the Data Exchange Web Service technical specifications. The technical specifications are updated periodically to reflect enhancements to the Data Exchange system and are on the Data Exchange website.

10.5.2 Bulk File Upload

Organisations with their own client management software systems capable of creating and exporting XML files can continue using this software to collect and transfer their performance data. Organisations will need to make a one-off adjustment (or ‘enhancement’) to their application in accordance with the Data Exchange bulk upload technical specifications. The technical specifications are updated periodically to reflect enhancements to the Data Exchange system and are on the Data Exchange website.
### 10.5.3 Free web-based portal

Organisations can use the Data Exchange web-based portal to manually input their data. Once saved in the portal, data is automatically submitted to the Data Exchange. The web-based portal can be used like a basic case management system, although it only allows for data that is relevant to performance reporting. This option is available for organisations who do not have their own proprietary client management software, or for those whose systems cannot accommodate the requirements to submit data through system-to-system transfers or bulk file upload.

The Data Exchange web-based portal collects the data requirements set out in this document and is available to all organisations funded to deliver in-scop program activities.

Organisations that already have their own case/client management system and submit their data by system-to-system transfers or bulk upload can access the web-based portal to use the Data Exchange functionality. For example, organisations who report information (consistent with the priority requirements) via a system-to-system transfer or bulk upload, may also use the web-based portal to record SCORE information about changes to their client's circumstances, goals and outcomes (consistent with the extended data items in the partnership approach). This approach is useful where the functionality for recording and reporting the extended data items is not available within an organisation's existing client management system.

Organisations who choose to report using both their client management systems (i.e. via a system-to-system transfer or bulk upload) and the web-based portal are able to view the records of their clients from the web-based portal to monitor and manage the services they provide to these clients.

### 10.6 Data Exchange user responsibilities

Every user of the Data Exchange is bound by the *Privacy Act 1988* and must ensure they meet these requirements at all times. Data Exchange users must ensure they only access records where a genuine need exists. Additionally, all Data Exchange users must ensure the data entered into the Data Exchange is accurate and up to date, to the best of their knowledge.

It is the organisation's responsibility to correct or delete any incorrect data. Organisations should notify the department as soon as practical if they become aware of any security or privacy breaches.

Organisations using the Data Exchange like a client management system also have an additional obligation to follow the protocols for providing notification and obtaining consent.

Data captured in the Data Exchange will only be used for the purpose for which it was captured. Data Exchange data will not be provided to other parties in Australia or elsewhere in the world for any other purpose.

### 10.7 Organisations no longer reporting via the Data Exchange

Organisations that report performance data in the Data Exchange are able to receive self-service reports on the data submitted for that period. They will not be able to enter any additional data for a period that has closed or for any periods where they do not have an active funding agreement.

If an organisation is continuing to report on other active activities in the Data Exchange they will have access to Data Exchange reports for all activities they are funded to deliver. Organisations retain access to the Data Exchange portal and self-service reports for at least one full reporting period (six months) after their last activity has ceased.

Where an organisation is unable to access the Data Exchange, this could be because their access to the portal is removed, or the organisation no longer has a user ID, or is experiencing other access restrictions related to AUSkey. In this situation, the organisation can request a copy of the report from the relevant Funding Arrangement Manager or the Data Exchange Helpdesk upon confirmation that the organisation:
has authority to receive this report as a former funding recipient
reported in the Data Exchange for the requested report period.

10.8 Training materials and help

Users of the Data Exchange web-based portal can access self-guided training material on the Data Exchange website.

Task cards
Task cards take users step-by-step through the processes required to create and manage records in the Data Exchange web-based portal.

e-Learning modules
Users of the Data Exchange can also access a suite of training videos known as e-Learning modules. These videos are on the Data Exchange website.

The Data Exchange Helpdesk
The Helpdesk is available to provide technical help to users of the Data Exchange.

You can contact the Data Exchange Helpdesk by email to dssdataexchange.helpdesk@dss.gov.au or phone 1800 020 283 between 8:30am – 5:30pm (AEST/AEDT), Monday to Friday.
## 11 List of data values

### Table 7. Priority requirements: client level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Client ID</td>
<td>4.4.2</td>
<td>Free text limit of 50 characters. If left blank a system generated number is assigned in the web-based portal, beginning at 001.</td>
</tr>
<tr>
<td>Given name *</td>
<td>5.1.1</td>
<td>Free text limit of 30 characters</td>
</tr>
<tr>
<td>Family name *</td>
<td>5.1.1</td>
<td>Free text limit of 30 characters</td>
</tr>
<tr>
<td>Date of birth *</td>
<td>5.1.2</td>
<td>Date format of dd/mm/yyyy</td>
</tr>
<tr>
<td>Estimated date of birth *</td>
<td>5.1.2</td>
<td>Tick box</td>
</tr>
<tr>
<td>Pseudonym used</td>
<td>5.1.1</td>
<td>Tick box</td>
</tr>
<tr>
<td>Gender *</td>
<td>5.1.3</td>
<td>Male</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Female</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Intersex/indeterminate</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Not stated or inadequately described</td>
</tr>
<tr>
<td>Residential address</td>
<td>5.1.4</td>
<td>Residential address line 1 (optional)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Address line 2 (optional)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Suburb (mandatory)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>State (mandatory)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Post code (limit of 4 digits) (mandatory)</td>
</tr>
<tr>
<td>Indigenous status</td>
<td>5.1.6</td>
<td>No</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Aboriginal</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Torres Strait Islander</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Aboriginal and Torres Strait Islander</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Not stated/inadequately described</td>
</tr>
<tr>
<td>Cultural and Linguistic Diversity: Country of Birth</td>
<td>5.1.7</td>
<td>Drop-down list of values based on the Australian Bureau of Statistics Standard Australian Classification of Countries (SACC), 2016</td>
</tr>
<tr>
<td>Cultural and Linguistic Diversity: Main language spoken at home</td>
<td>5.1.7</td>
<td>Drop-down list of values based on the Australian Bureau of Statistics Australian Standard Classification of Languages (ASCL), 2016</td>
</tr>
<tr>
<td>Data Field</td>
<td>Protocols Section</td>
<td>Field Values</td>
</tr>
<tr>
<td>------------------------------------------------</td>
<td>-------------------</td>
<td>-----------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Disability, impairment or condition indicator</td>
<td>5.1.8</td>
<td>Intellectual/learning, Psychiatric, Sensory/speech, Physical/diverse, None (no disability), Not stated/inadequately described</td>
</tr>
<tr>
<td>Consent to have personal information stored in the web-based portal</td>
<td>4.2.3</td>
<td>Tick box</td>
</tr>
<tr>
<td>Consent to participate in follow up research, surveys and evaluation</td>
<td>4.2.5</td>
<td>Tick box</td>
</tr>
</tbody>
</table>

*These fields generate an AIHW Statistical Linkage Key (SLK) 581 algorithm.

**Table 8. Priority requirements: case level data**

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case ID</td>
<td>5.2.1</td>
<td>Free text limit of 50 characters. If left blank a system generated number is assigned.</td>
</tr>
<tr>
<td>Outlet</td>
<td>3.5</td>
<td>In the web-based portal: to be selected from a list of options in the drop-down.</td>
</tr>
<tr>
<td>Program Activity</td>
<td>5.2.1</td>
<td>In the web-based portal: to be selected from a list of options in the drop-down. The drop-down will only display program activities that are assigned to the outlet selected.</td>
</tr>
<tr>
<td>Unidentified client count</td>
<td>2.6</td>
<td>Free text number only with limit of 999</td>
</tr>
<tr>
<td>Attach clients</td>
<td>2.2</td>
<td>In the web-based portal: to be selected from a list of options in the drop-down. The drop-down provides a mechanism to associate one or more client records to the case.</td>
</tr>
</tbody>
</table>
Table 9. Priority requirements: session level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Session ID</td>
<td>5.2.2</td>
<td>Free text limit of 50 characters. If left blank a system generated number is assigned.</td>
</tr>
<tr>
<td>Session date</td>
<td>5.2.2</td>
<td>Date format of dd/mm/yyyy</td>
</tr>
<tr>
<td>Unidentified clients attending this session (optional)</td>
<td>5.2.2</td>
<td>Number field. The default value is 0, maximum 999 (however cannot exceed the value specified at the Case level).</td>
</tr>
<tr>
<td>Client attendance</td>
<td>5.2.2</td>
<td>Record for each case clients present at the session.</td>
</tr>
<tr>
<td>Service type</td>
<td>5.2.2</td>
<td>The number and variety of service types will depend on the program activity selected. The full list of values is in the Service Type Matrix on the Data Exchange website.</td>
</tr>
</tbody>
</table>

Table 10. Commonwealth Home Support Programme: client level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
</table>
| Accommodation setting             | 5.3.1             | Boarding house  
Crisis, emergency or transition  
Independent living unit  
Indigenous community/settlement  
Institutional setting (i.e. residential aged care, hospital)  
Private residence—client or family owned/purchasing  
Private residence—private rental  
Private residence—public rental  
Public shelter  
Supported accommodation  
Other  
Not stated |
<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Living arrangements</td>
<td>5.3.1</td>
<td>Single (person living alone)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Sole parent with dependant(s)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Couple</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Couple with dependant(s)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Group (related adults)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Group (unrelated adults)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Homeless/no household</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Not stated or inadequately described</td>
</tr>
<tr>
<td>DVA card status</td>
<td>5.3.1</td>
<td>DVA Gold Card</td>
</tr>
<tr>
<td></td>
<td></td>
<td>DVA White Card</td>
</tr>
<tr>
<td></td>
<td></td>
<td>DVA Orange Card or other</td>
</tr>
<tr>
<td></td>
<td></td>
<td>No DVA entitlement</td>
</tr>
<tr>
<td>Existence of Carer</td>
<td>5.3.1</td>
<td>Yes</td>
</tr>
<tr>
<td></td>
<td></td>
<td>No</td>
</tr>
<tr>
<td>Client exit reason</td>
<td>5.3.1</td>
<td>Client no longer requires assistance</td>
</tr>
<tr>
<td>(program specific only)</td>
<td></td>
<td>Service unable to provide assistance</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Client now requires higher level of aged care</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Client has moved out of area</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Client terminated the service</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Client died</td>
</tr>
<tr>
<td></td>
<td></td>
<td>None of the above</td>
</tr>
</tbody>
</table>

Table 11. Commonwealth Home Support Programme: session level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Amount of assistance</td>
<td>5.3.1</td>
<td>The information required for this field will depend on the service type</td>
</tr>
<tr>
<td>provided</td>
<td></td>
<td>selected. Go to the program specific guidance documents on the website to</td>
</tr>
<tr>
<td></td>
<td></td>
<td>determine which fields apply to each service type:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Hours/minutes</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Quantity</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Cost</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Type</td>
</tr>
<tr>
<td>Fees charged</td>
<td>5.3.1</td>
<td>Number field (whole dollars only) appears where applicable</td>
</tr>
</tbody>
</table>
Table 12. Family Law activities: case/session level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fees charged</td>
<td>5.3.2</td>
<td>Number field (whole dollars only) appears where applicable</td>
</tr>
<tr>
<td>Was a parenting agreement reached?</td>
<td>5.3.2</td>
<td>Appears, where applicable when a case is created. The values are:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Full</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Partial</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Not reached</td>
</tr>
<tr>
<td>Date of parenting agreement</td>
<td>5.3.2</td>
<td>Date format of dd/mm/yyyy</td>
</tr>
<tr>
<td>Did a legal practitioner assist with formalising agreement?</td>
<td>5.3.2</td>
<td>Yes</td>
</tr>
<tr>
<td></td>
<td></td>
<td>No</td>
</tr>
<tr>
<td>Section 60(I) certificate type</td>
<td>5.3.2</td>
<td>Attended—genuine effort</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Attended—not genuine effort</td>
</tr>
<tr>
<td></td>
<td></td>
<td>FDR began—considered inappropriate to continue</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Matter inappropriate for resolution</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Not held due to refusal or failure of other person to attend</td>
</tr>
<tr>
<td>Date issued (Section 60(I) Certificate)</td>
<td>5.3.2</td>
<td>Date format of dd/mm/yyyy</td>
</tr>
</tbody>
</table>

Table 13. Financial Wellbeing and Capability: session level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Which money management course was delivered?</td>
<td>5.3.3</td>
<td>This question only appears if an applicable activity and service type is selected when the case and session is created. Applicable values are:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 1 - Making Money Last Until Payday</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 2 - Planning For the Future</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 3 - How Can Banks Help</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 4 - Internet and Phone Banking</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 5 - Credit Can Be a Hazard</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 6 - Money Loans Sharks and Traps</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 7 - A Roof Overhead – Home Ownership</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 8 - A Roof Overhead Tenancy</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Workshop 9 - Managing Paperwork</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Other workshop</td>
</tr>
</tbody>
</table>
### Table 14. National Disability Advocacy Program: session level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Topic</td>
<td>5.3.5</td>
<td>Abuse/Neglect/Violence</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Access to non NDIS service</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Child Protection</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Community Inclusion—Social/Family</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Disability services complaints</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Discrimination/rights</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Education</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Employment</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Equipment/aids</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Finances</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Government payments</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Health/ Mental Health</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Housing/Homelessness</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Legal/Access to Justice</td>
</tr>
<tr>
<td></td>
<td></td>
<td>NDIS—Internal Review</td>
</tr>
<tr>
<td></td>
<td></td>
<td>NDIS—Access/Planning</td>
</tr>
<tr>
<td></td>
<td></td>
<td>NDIS—Support implementing plan/Accessing services</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Other</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Physical access</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Transport</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Vulnerable/isolated</td>
</tr>
</tbody>
</table>
Table 15. Partnership approach: client level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Homeless indicator</strong></td>
<td>6.10</td>
<td>Yes</td>
</tr>
<tr>
<td></td>
<td></td>
<td>No</td>
</tr>
<tr>
<td></td>
<td></td>
<td>At Risk</td>
</tr>
<tr>
<td><strong>Household composition</strong></td>
<td>6.5</td>
<td>Single (person living alone)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Sole parent with dependant(s)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Couple</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Couple with dependant(s)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Group (related adults)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Group (unrelated adults)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Homeless/No household</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Not stated or inadequately described</td>
</tr>
<tr>
<td><strong>Highest level of education/qualification</strong></td>
<td>6.6</td>
<td>Pre-primary education</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Primary education</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Secondary education</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Certificate level</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Advanced diploma and diploma level</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Bachelor degree level</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Graduate diploma and graduate certificate level</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Postgraduate degree level</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Other education</td>
</tr>
<tr>
<td>Data Field</td>
<td>Protocols Section</td>
<td>Field Values</td>
</tr>
<tr>
<td>--------------------------</td>
<td>-------------------</td>
<td>-------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Employment status</td>
<td>6.7</td>
<td>Paid work full-time&lt;br&gt;Paid work part-time&lt;br&gt;Unpaid work (includes volunteering)&lt;br&gt;Not working and not looking for work&lt;br&gt;Unemployed (not working but looking for work)&lt;br&gt;Studying full-time&lt;br&gt;Studying part-time&lt;br&gt;Caring&lt;br&gt;Parenting</td>
</tr>
<tr>
<td>Main source of income</td>
<td>6.8.1</td>
<td>Nil income&lt;br&gt;Employee salary/wages&lt;br&gt;Other income including superannuation and investments&lt;br&gt;Self-employed (unincorporated business income)&lt;br&gt;Government payments/pensions/allowances&lt;br&gt;Not stated/Inadequately described</td>
</tr>
<tr>
<td>Income frequency</td>
<td>6.8.2</td>
<td>Weekly&lt;br&gt;Fortnightly&lt;br&gt;Monthly&lt;br&gt;Annually</td>
</tr>
<tr>
<td>Approximate gross income</td>
<td>6.8.2</td>
<td>Number field (whole dollars only)</td>
</tr>
<tr>
<td>Month of first arrival in Australia</td>
<td>6.9</td>
<td>Drop-down menu of twelve month calendar year</td>
</tr>
<tr>
<td>Year of first arrival in Australia</td>
<td>6.9</td>
<td>Drop-down menu of year in chronological order</td>
</tr>
<tr>
<td>Visa type</td>
<td>6.9</td>
<td>Humanitarian&lt;br&gt;Family&lt;br&gt;Skilled&lt;br&gt;Other</td>
</tr>
<tr>
<td>Ancestry</td>
<td>6.9</td>
<td>Select from the list of values which is based on the Australian Bureau of Statistics <em>Australian Standard Classification of Cultural and Ethnic Groups (ASCCEG), 2016</em></td>
</tr>
<tr>
<td>Is client a carer</td>
<td>6.12</td>
<td>Yes&lt;br&gt;No</td>
</tr>
</tbody>
</table>
### Table 16. Partnership approach: case level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Attendance profile</td>
<td>6.11</td>
<td>Family, Community event, Peer support group, Couple, Cohabitants</td>
</tr>
<tr>
<td>Reason for seeking</td>
<td>6.2</td>
<td>Physical health, Mental health, wellbeing and self-care, Personal and family safety, Age-appropriate development, Community participation and networks, Family functioning, Financial resilience, Employment, Education and skills training, Material wellbeing and basic necessities, Housing</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>NDIS eligibility</td>
<td>6.13</td>
<td>NDIS in-progress access request, NDIS eligible, NDIS ineligible</td>
</tr>
<tr>
<td>Data Field</td>
<td>Protocols Section</td>
<td>Field Values</td>
</tr>
<tr>
<td>------------------</td>
<td>-------------------</td>
<td>-----------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Referral source</td>
<td>6.3</td>
<td>Health agency</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Community services agency</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Educational agency</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Internal</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Legal agency</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Employment/job placement agency</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Centrelink/Department of Human Services (DHS)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Other Agency</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Self</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Family</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Friends</td>
</tr>
<tr>
<td></td>
<td></td>
<td>General Medical Practitioner</td>
</tr>
<tr>
<td></td>
<td></td>
<td>My Aged Care Gateway</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Linkages Package</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Continuity of Support (CoS) Programme</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Humanitarian Settlement Program</td>
</tr>
<tr>
<td></td>
<td></td>
<td>LAC Referral</td>
</tr>
<tr>
<td></td>
<td></td>
<td>NDIS referral</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Other party</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Not stated/inadequately described</td>
</tr>
</tbody>
</table>

Table 17. Partnership approach: session level data

<table>
<thead>
<tr>
<th>Data Field</th>
<th>Protocols Section</th>
<th>Field Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Service setting</td>
<td>6.14</td>
<td>Organisation outlet/office</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Clients residence</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Community venue</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Partner organisation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Telephone</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Digital</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Healthcare facility</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Education facility</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Justice facility</td>
</tr>
<tr>
<td>Interpreter present</td>
<td>6.15</td>
<td>Yes</td>
</tr>
<tr>
<td></td>
<td></td>
<td>No</td>
</tr>
<tr>
<td>Data Field</td>
<td>Protocols Section</td>
<td>Field Values</td>
</tr>
<tr>
<td>----------------------------</td>
<td>-------------------</td>
<td>----------------------------------------------------------------------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Referral type</td>
<td>6.4</td>
<td>Internal—made to another service offered within the same organisation</td>
</tr>
<tr>
<td></td>
<td></td>
<td>External—made to a service that is provided by a different organisation</td>
</tr>
<tr>
<td>Referral purpose</td>
<td>6.4</td>
<td>Physical health</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Mental health, wellbeing &amp; self-care</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Personal and family safety</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Age-appropriate development</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Community participation &amp; networks</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Financial Resilience</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Family functioning</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Employment</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Education and skills training</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Material wellbeing and basic necessities</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Housing</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Support to caring role</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Other</td>
</tr>
<tr>
<td>Circumstance SCORE domains</td>
<td>7.2.1</td>
<td>Physical health</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Mental health, wellbeing and self-care</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Personal and family safety</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Age-appropriate development</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Community participation and networks</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Family functioning</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Financial resilience</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Employment</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Education and skills training</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Material wellbeing and basic necessities</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Housing</td>
</tr>
<tr>
<td>Goals SCORE domains</td>
<td>7.3.1</td>
<td>Changed knowledge and access to information</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Changed skills</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Changed behaviours</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Empowerment, choice and control to make own decisions</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Engagement with relevant support services</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Changed impact of immediate crisis</td>
</tr>
<tr>
<td>Satisfaction SCORE</td>
<td>7.4.1</td>
<td>I am satisfied with the services I have received</td>
</tr>
<tr>
<td>domains</td>
<td></td>
<td>The service listened to me and understood my issues</td>
</tr>
<tr>
<td></td>
<td></td>
<td>I am better able to deal with issues that I sought help with</td>
</tr>
<tr>
<td>Data Field</td>
<td>Protocols Section</td>
<td>Field Values</td>
</tr>
<tr>
<td>----------------------------</td>
<td>-------------------</td>
<td>-------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Community SCORE domains</td>
<td>7.5.1</td>
<td>Community infrastructure and networks</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Organisational knowledge, skills and practices</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Group/community knowledge, skills, attitudes and behaviours</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Social cohesion</td>
</tr>
<tr>
<td>Assessed by</td>
<td>7.1</td>
<td>SCORE directly – client</td>
</tr>
<tr>
<td></td>
<td></td>
<td>SCORE directly – practitioner</td>
</tr>
<tr>
<td></td>
<td></td>
<td>SCORE directly – joint</td>
</tr>
<tr>
<td></td>
<td></td>
<td>SCORE directly – support person</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Validated outcomes tool – client</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Validated outcomes tool – practitioner</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Validated outcomes tool – joint</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Validated outcomes tool – support person</td>
</tr>
</tbody>
</table>
12 Version history

Version 6, October 2019

The department released version 6 of the protocols following a review of readability and accessibility. All sections were changed to improve reading ease.

The section on client surveys was clarified to reflect that the department will no longer be implementing a generalised outcomes survey to all Data Exchange clients. Instead, program specific surveys will be implemented where required.

Version 5, November 2018

The department released version 5 of the protocols in October 2018 to reflect the August 2018 system enhancements. Details of the August 2018 enhancements are on the Data Exchange website. The changes are:

- **Various sections** - ‘service provider’ replaced with ‘organisation’. This is a better reflection of who is reporting into the Data Exchange.
- **Section 5** - addition of program specific fields for Career Pathways Pilot for Humanitarian Entrants and the National Disability Advocacy Program
- **Section 6** - updated references to the partnership approach
- **Section 7** - updated naming, definition and scope of Circumstances outcome domains
- **Section 8** - updated client survey pilot references
- **Section 9** - information on the handshake agreement
- **Section 11** - updated data values made as part of the August 2018 enhancements

Version 4, March 2017

This review was conducted in the context of the Data Exchange expansion to other agencies and jurisdictions, and the creation of the Community Grants Hub. The changes are:

- **Section 6** - addition of the partnership approach as a requirement of some program funding agreements
- **Section 8** - updated with client survey and changed section name and numbering
- **Section 9** - updated with reports information and changed section name and numbering
- **Section 10** - updated program specific fields for Career Pathways Pilot for Humanitarian Entrants and the National Disability Advocacy Program. Changed section name and numbering
- **Section 11** - addition of new Data Values
- **Section 12** - addition of version history section (previously Attachments 1 & 2)

Version 3, October 2015

The department released version 3 of the protocols in October 2015 to include enhancements made to the system for the Commonwealth Homes Support Programme activity and clarify protocols relating to protecting the client’s privacy and personal information.

- **Section 2** - Added an example for who is a client for the Commonwealth Home Support Programme (CHSP)
- **Section 3** - Included definition of outlet
- **Section 4** - Consolidated information about personal information; Limited the content to the department’s privacy obligations; Updated the consent and notification protocols and arrangements for follow-up research.
- **Section 5** - Clarified how to report the client’s residential address; introduced data items for CHSP.
- **Section 6** - Added homeless/no household value in the extended data items for household composition; introduced client exit reason for CHSP cases.
- **Section 7** - Added the Translation Matrix for SCORE; moved in-scope activities and service types into Appendix A—Service Type Matrix.
- **Section 8** - Expanded client management functions of the Data Exchange; Clarified the Statistical Linkage Key.

Version 2, March 2015

The department released version 2 of the protocols in March 2015 to align the Data Exchange technical specifications (December 2014) and the Data Exchange web-based portal (February 2015).

The original version of this document was published in August 2014.