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| Requirements of SCIg approval for health services |

**Health services must take full accountability for use and management of SCIg product.**

**Supply of product**

Managed via BloodSTAR. Maximum 2 months’ supply

**Equipment and facilities**

Provision of all necessary equipment and consumables to administer SCIg

Further information [SCIg program, tools and resources (health.vic.gov.au)](https://www.health.vic.gov.au/patient-care/scig-program-tools-and-resources)

**Regular review**

Assess clinical benefit of SCIg. Patients can record SCIg product use and any adverse reactions to aid the review process.

**Clinical oversight**

Supervision by specialist and ongoing assessment of clinical benefit for patients assessed as suitable for SCIg. Ongoing clinical oversight and support for participating patients

**Complete Hospital Acknowledgement Form.**

National Subcutaneous Immunoglobulin

https://[www.blood.gov.au/SCIg](http://www.blood.gov.au/SCIg)

Submit to National Blood Authority for approval:

Attention: Ig Governance Fax: (02) 6151 5235

Email: iggovernance@blood.gov.au

**Education and training**

Must be provided for staff and patients to ensure appropriate management and use of SCIg.

**Governance**

**Reporting unused, discarded, spoilt/broken product**

Reported via BloodNet.

**Quality assurance**

Policies and procedures for SCIg management.

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