

# Advancing quality in IV and SubQ immunoglobulin therapy



### Your partner in patient care

At Advanced Infusion Care, a division of AIS Healthcare, we apply a team approach to intravenous (IV) and subcutaneous (SubQ) immunoglobulin therapy patient care.

Collaboration between physicians and the nationally accredited AIC home infusion team ensures the consistent delivery of patient-specific, specialized in-home infusion services to patients across the country, helping to improve outcomes—and lives.

#### **DOING MORE OF WHAT MATTERS**

#### ✓ A team approach to infusion care

Every AIC patient has 24/7 access to a dedicated team of experienced clinical and support staff, including Clinical Pharmacists, Infusion Nurse Specialists, Patient Care Managers, and Intake Managers.

#### ✓ Nationally recognized quality

Dually accredited by URAC and the Accreditation Commission for Health Care (ACHC), AIC is a member of the Immunoglobulin National Society (IgNS) and is regularly inspected by National and State Boards of Pharmacy.

#### ✓ Billing and reimbursement made easy

We work with patients, our large network of payers, and your office to secure needed authorizations and complete clinical paperwork as well as offer financial assistance to qualifying patients.

#### ✓ Supply to meet your needs

We have long-term agreements with multiple suppliers to provide you and your patients with the right Ig infusion therapy.



## How to refer patients to AIC

At AIC, we want to make patient onboarding as easy as possible. Use the documentation checklists and associated ICD-10 diagnosis codes below to send us the required patient information—we'll take it from there.

	ATION NEEDED FOR PRIMARY IMMUNE (PI) NCY DISEASE REFERRALS:	INFORMATION NEEDED FOR NEUROLOGY THERAPY REFERRALS:							
<ul> <li>Copy</li> <li>Presci</li> <li>H&amp;P (i of infe</li> <li>Serum</li> <li>Ig1, Ig2</li> <li>Recer</li> <li>Vaccir</li> </ul>	of patient's insurance card ription (including dose and frequency) ncluding supporting documentation ction history) n immunoglobulin levels (including IgA, IgG, and IgM) 2, Ig3, and Ig4 subclass report (if available) nt BUN and creatinine results ne challenge test results and titer values alar Access Device (VAD) report, if applicable	<ul> <li>Patient demographic sheet</li> <li>Copy of patient's insurance card</li> <li>Prescription (including dose and frequency)</li> <li>H&amp;P</li> <li>Recent BUN and creatinine results</li> <li>Nerve conduction velocity study</li> <li>Lumbar puncture showing CSF protein levels</li> <li>Diagnostic studies: nerve conduction studies/EMG/muscle biopsy/CK levels</li> <li>Vascular Access Device (VAD) report, if applicable</li> </ul>							
	D-10 codes for PI therapy*: DE DESCRIPTION		CD-10 codes for neurology therapy*: DE DESCRIPTION						
D80.0†	Hereditary hypogammaglobulinemia	D89.8	Disorder involving the immune mechanism, unspecified						
D80.1	Nonfamilial hypogammaglobulinemia	G25.82	Stiff-man syndrome						
D80.2 <sup>‡</sup>	Selective deficiency of IgA	G35	Multiple sclerosis (RRMS)						
D80.3‡	Selective deficiency of IgG subclasses	G60.9	Hereditary and idiopathic neuropathy, unspecified						
D80.5‡	Immunodeficiency with increased IgM	G61	Inflammatory polyneuropathy						
D80.6‡	Antibody deficiency with near-normal immunoglobulins	G61.0	Guillain-Barre syndrome						
D81.0 <sup>†</sup>	SCID with reticular dysgenesis	G61.81	Chronic inflammatory demyelinating polyneuritis						
D81.1 <sup>†</sup>	SCID with low T- and B-cell numbers	G61.82	Multifocal motor neuropathy						
D81.2 <sup>†</sup>	SCID with low or normal B-cell numbers	G61.9	Inflammatory polyneuropathy, unspecified						
D81.6†	Major histocompatibility complex class I deficiency	G70.0	Myasthenia gravis and other myoneural disorders						
D81.7 <sup>†</sup>	Major histocompatibility complex class II deficiency	G70.01	Myasthenia gravis with (acute) exacerbation						
D81.89 <sup>†</sup>	Other combined immunodeficiencies	G70.80	Lambert-Eaton syndrome, unspecified						
D81.9†	Combined immunodeficiency, unspecified	G72.41	Inclusion body myositis						
D82.0†	Wiskott-Aldrich syndrome	G72.49	Other inflammatory and immune myopathies						
D82.9	Immunodeficiency associated with major defect, unspecified	G73.1	Lambert-Eaton syndrome in neoplastic disease						
D83.0 <sup>†</sup>	CVID with predominant abnormalities of B-cell numbers and function	M34.82	Scleroderma						
D83.1‡	CVID with predominant immunoregulatory T-cell disorders	M32.19	Systemic lupus erythematosus						
D83.2 <sup>†</sup>	CVID with autoantibodies to B- or T-cells	M33.1	Dermatomyositis						
D83.8†	Other common variable immunodeficiencies	M33.2	Polymyositis						
D83.9†	CVID, unspecified								

<sup>\*</sup>These ICD-10 codes fall under the disease states listed in Jolles S et al. Clinical uses of intravenous immunoglobulin. Clin Exp Immunol. 2005;142(1):1-11. doi:10.1111/j.1365-2249.2005.02834

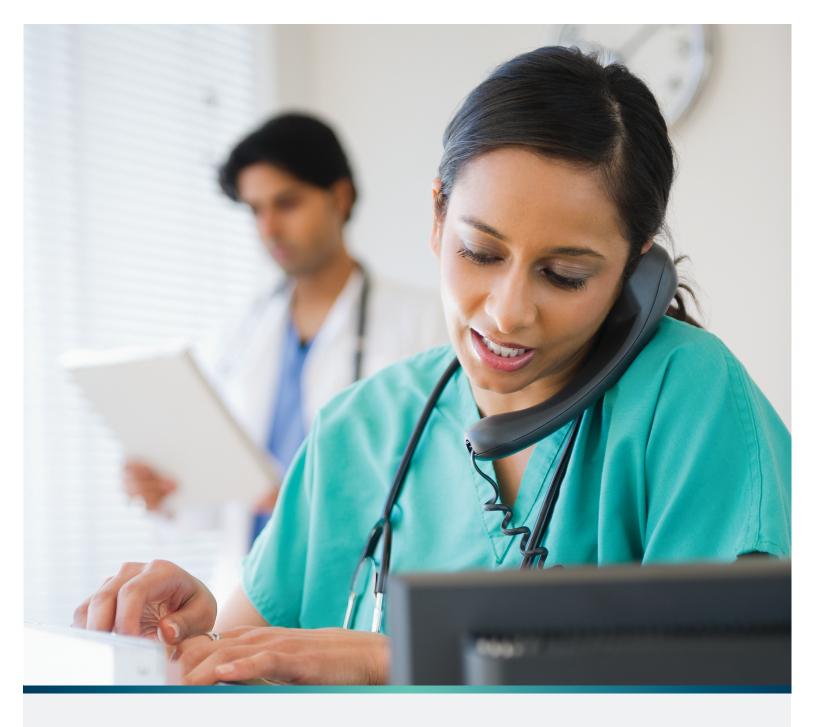
†These ICD-10 codes reflect diagnoses that are payable for Ig home infusion under Medicare Part B as published in IDF: SCID Compass. More PI diagnoses covered for home Ig replacement therapy under Medicare Part B. July 2019. https://primaryimmune.org/scid-compass/news/more-pi-diagnoses-covered-home-ig-replacement-therapy-under-medicare-part-b.

‡These ICD-10 codes were added per CMS guidelines, effective August 2019, as published in IDF: SCID Compass. See reference directly above.

## Discover our selection of immunoglobulin products

Learn more about the IV and SubQ immunoglobulin therapy products that we offer. Want selection or dosing guidance? Our clinical pharmacists are ready to help you pick the products that meet the health conditions and restrictions of your patients.

	ASCENIV™	BIVIGAM°	GAMM	APLEX*	HIZENTRA®	PRIVIGEN°	GAMUNEX*-C XEMBIFY*		XEMBIFY*	GAMMAKED™		CUTAQUIG*	OCTAGAM <sup>®</sup>	PANZYGA*	CUVITRU**	GAMMAGARD LIQUID*	GAMMAGARD® S/D	HYQVIA°
Manufacturer	ADMA Bio	ologics Inc.	Bio Products Laboratory		CSL Behring		Grifols		Kedrion		Octapharma		Pfizer		Takeda			
Indications	PI	PI	PI,ITP		PI, CIDP	PI, ITP	IV: PI, SubQ: ITP, PI CIDP		PI	IV: PI, ITP, CIDP	SubQ: PI	PI	5%:PI 10%: ITP dermatomyositis	PI,ITP	PI (patients >2 yrs of age)	IV:PI, SubQ: MMN PI	PI, ITP, B-cell CLL, Kawasaki disease	PI
Form	Liquid	Liquid	Liquid		Liquid	Liquid	Liquid Liquid		Liquid		Liquid	Liquid	Liquid	Liquid	Liquid	Lyophilized	Liquid	
Shelf life and storage requirements	Refrigerate at 2-8°C (36-46°F). Do not freeze or heat. Do not use after expiration date	Stored until expiration date on vial packaging at 2-8°C (36-46°F)	36 months (room temperature storage)		30 months (room temperature storage)	36 months (room temperature storage)	36 months (refrigerated at 36-46°F) 6 months (room temperature storage not exceeding 77°F)		36 months		24 months (refrigerated) 6 months (room temperature storage not exceeding 77°F)	24 months	24 months (refrigerated up to 46°F) 9 months (room temperature storage not exceeding 77°F)	12 months room temperature (do not exceed 25°C/77°F)	36 months (refrigerated) 24 months (room temperature storage not exceeding 77°F)	24 months (room temperature storage)	36 months (refrigerated at 36-46°F) 3 months (room temperature storage not exceeding 77°F)	
Reconstitution time	None (liquid solution)	None (liquid solution)	None (liquid solut	ion)	None (liquid solution)	None (liquid solution)	None (liquid soli	ution)	None (liquid solution)	None (liquid solution)		None (liquid solution)	None (liquid solution)	None (liquid solution)	None (liquid formulation)	None (liquid solution)	n/a	None (liquid solution)
Available concentration	10%	10%	5%	10%	20%	10%	10% 20%		20%	10%		16.5%	5% 10%	10%	20%	10%	5%	10%
Maximum recommended infusion rate	Up to 0.08 mL/kg per minute	Up to 6 mL/kg/ minute	4.8 mL/kg per ho	ur	Up to 25 mL/ hour per site (50 mL/hour for all sites combined)	4.8 mL/kg per hour	4.8 mL/ kg per hour	20 mL/ per hour	25 mL/hour per site	4.8 mL/ kg per hour	20 mL/ per hour	Up to 100 mL/ hour at all sites combined	≤4.2 mL/kg per hour	0.01 mL/kg per minute	First 2 infusions: 10-20 mL/hr/site All subsequent infusions: up to 60 mL/ hr/site	5 mL/kg ≥40 kg BW:  per hour 30 mL/site at  20-30 mL/ hour per site  ≤40 kg BW:  20 mL/site at  15-20 mL/hour per site	4 mL/kg per hour	<40 kg: ≤300 mL per injection site ≥40 kg: ≤600 mL per injection site
Time to infuse 35 g	Varies based on volume and tolerability	Varies based on volume and tolerability	2 hrs, 40 min for a 70-kg person, if infused according to PI	1 hr, 53 min for a 70-kg person, if infused according to PI	Varies based on volume and tolerability	Varies based on patient tolerability	Varies based on administration method		Varies based on volume and tolerability	Varies based on administration method		Varies based on patient tolerability	2 hours, 30 minutes	Varies based on patient tolerability	Varies based on patient tolerability	Varies based on patient tolerability	Varies based on patient tolerability	Varies based on patient tolerability
Sugar content	Contains no sucrose	Contains no sucrose/glucose/ maltose	5% D-sorbitol (polyol)	None	None	None	None		None	None		79 mg/mL (maltose)	10 mg/mL (maltose)	None	No added sugar	No added sugars	20 mg/mL (glucose)	No added sugars
Sodium content	0.100-0.140 M (sodium chloride)	0.100-0.140 M (sodium chloride)	30-50 mmol/L	<30 mM	Trace amounts (≤10 mmol/L)	Trace amounts	Trace amounts		Trace amounts	Trace amounts		≤30 mmol/L	≤30 mmol/L	Trace amounts	No added sodium	No added sodium	8.5 mg/mL (sodium chloride)	8.5 mg/mL in HYQVIA (none in immunoglobulin)
Osmolarity/ osmolality	240-310 mOsm/kg	454-472 mOsm/kg	460-500 mOsm/kg	~280 mOsm/kg	380 mOsm/kg	Isotonic (380 mOsm/kg)	258 mOsm/kg		280-404 mOsm/kg	258 mOsm/kg		310-380 mOsm/kg	310-380 mOsm/kg	240-310 mOsm/kg	280-292 mOsm/kg	240-300 mOsm/kg	636 mOsm/kg	240-300 mOsm/kg
рН	4.0-4.6	4.0-4.6	4.6-5.1	4.9-5.2	4.6-5.2	4.8	4.0-4.5		4.1-4.8	4.0-4.5		5.0-5.5	5.1-6.0	4.5-5.0	4.6-5.1	4.6-5.1	6.8 ± 0.4	4.6-5.1
IgA content	≤200 µg/mL	Contains trace amounts of IgA	<4 mcg/mL (average)	<20 mcg/mL (specification value)	≤50 mcg/mL	≤25 mcg/mL	46 μg/mL		IgA <0.07 mg/ mL	46 μg/mL		≤0.6 mg/mL	100 μg/mL	100 µg/mL (average)	80 µg/mL	37 μg/mL	<1µg/mL	37 μg/mL
Approved method of administration	IV	IV	IV		SubQ	IV	IV	SubQ	SubQ	IV	SubQ	SubQ	IV	IV	SubQ	IV SubQ	IV	SubQ



## It's easy to get started

At AIC, we want to be your partner in patient care. To get your patients started on infusion therapy, simply complete a referral form—forms for both IV and SubQ therapies are on our website (advancedinfusioncare. com). Submit the referral form along with the patient's Vascular Access Device (VAD) report, if applicable. Once the referral is evaluated by a clinical review specialist, an AIC representative will be in touch.

To learn more about our in-home infusion services or the products we offer, please contact us.