

Broad Access to Immunoglobulin (IG) Therapies

As the largest independent infusion services provider in the U.S.¹, Option Care Health meets the comprehensive needs of autoimmune patients. Our extraordinary team of highly skilled professionals are committed to quality and exceptional patient care.

Extensive payer coverage

 $o 96\%^2$ coverage to insured lives

Clinical pharmacy appeals team, supporting all IG claims

- Patient assistance program expertise
 - o Our integrated system matches eligible patients with manufacturer copay assistance and foundation programs coupled with an expedited income verification process, reducing insurance approval time
- Flexible infusion therapy settings and the largest infusion therapy nursing network

o Patient's Home | Infusion Suite | 170+² nationwide locations

Immunoglobulin Therapy

Multiple brands of IG therapies for personalized care

One simple choice

for extraordinary infusion care

 Only national infusion provider in-network with every national health plan including their affiliates¹

o National | Regional | Medicare | Medicaid

 Treat a wide range of acute and chronic conditions through our comprehensive infusion portfolio

o Clinically managing 95,000+¹ IG infusions each year

• Transitioning patient's onto care quickly

o Dedicated patient intake reduces administrative burdens

• 93%² patient satisfaction rating

	Intravenous/Subcutaneous					
Product	Gammagard [®] Liquid	Gammaked™	Gamunex®-C			
Manufacturer	Takeda Pharmaceutical Company Limited 800.828.2088 takeda.com gammagard.com	Kedrion Biopharma, Inc. 855.353.7466 kedrion.us gammaked.com	Grifols 888.694.2686 grifols.com gamunex-c.com			
Indications	PI (adults & peds 2+ yrs.) MMN (adults, maintenance therapy) CIDP (adult)	PI (adults & peds 2+ yrs.) ITP (adults & children) CIDP (adults)	PI (adults & peds 2+ yrs.) ITP (adults & children) CIDP (adults)			
Contraindications	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy			
Form	Liquid	Liquid	Liquid			
Concentration	10%	10%	10%			
Sizes	1/2.5/5/10/20/30 g	1/2.5/5/10/20 g	1/2.5/5/10/20/40 g			
Virus removal method	Cohn-Oncley cold ethanol fractionation, cation/ anion exchange chromatography, solvent/deter- gent, nanofiltration, low pH incubation	Cold ethanol fractionation, caprylate precipita- tion/filtration, anion exchange chromatography, low pH incubation	Cold ethanol fractionation, caprylate precipita- tion/filtration, anion exchange chromatography, low pH incubation			
IgA content	37 μg/mL	46 μg/mL	46 μg/mL			
Stabilizer	Glycine	Glycine	Glycine			
Sugar content	None	None	None			
Sodium content	None	Unspecified	Unspecified			
рН	4.6-5.1	4.0-4.5	4.0-4.5			
Osmolality (mOsmol/kg)	240-300	258	258			
Half-life (days)	31-42	35.7	35.7			
Administration	 PI/MMN: IV: Begin at 0.8 mg/kg/min for first 30 minutes, if tolerated, gradually increase every 30 minutes to maximum infusion rate of 8 mg/kg/min as tolerated). PI: SubQ: Begin 1 week after last IVIG dose. (≥ 40kg) begin with 30 mL/site @ 20 mL/hr/site; maintenance 30 mL/site @ 15 mL/hr/ site; c< 40kg) begin with 20 mL/site @ 15-20 mL/hr/ site; maintenance 20 mL/site @ 15-20 mL/hr/ site. 	PI/ITP: IV: Begin at 1 mg/kg/min for first 30 minutes, if tolerated, gradually increase to maximum infusion rate of 8 mg/kg/min. CIDP: IV: Begin at 2 mg/kg/min for first 30 minutes, if tolerated, gradually increase to maximum infusion rate of 8 mg/kg/min. PI: SubQ: Begin 1 week after last IVIG dose. (adults) infuse @ 20 mL/hr/site; (children/adolescents ≥ 25kg) initially infuse @ 15 mL/hr/site; maintenance @ 20 mL/hr/site; (children/ adolescents < 25kg) infuse @ 10 mL/hr/site.	PI/ITP: IV: Begin at 1 mg/kg/min for first 30 minutes, if tolerated, gradually increase to maximum infusion rate of 8 mg/kg/min. CIDP: IV: Begin at 2 mg/kg/min for first 30 minutes, if tolerated, gradually increase to maximum infusion rate of 8 mg/kg/min. PI: SubQ: Begin 1 week after last IVIG dose. (adults) infuse @ 20 mL/hr/site; (children/adolescents ≥ 25kg) initially infuse @ 15 mL/hr/site; maintenance @ 20 mL/hr/site; (children/ adolescents < 25kg) infuse @ 10 mL/hr/site.			
Shelf-life/storage requirements	Refrigerated 2 to 8°C [36 to 46°F] for 36 months OR room temperature up to 25°C [77°F] for 24 months	Refrigerated 2 to 8°C [36 to 46°F] for 36 months OR room temperature up to 25°C [77°F] for 6 months	Refrigerated 2 to 8°C [36 to 46°F] for 36 months OR room temperature up to 25°C [77°F] for 6 months			

	Intravenous/Subcutaneous				
Product	Gammagard® Liquid	Gammaked™	Gamunex [®] -C		
Manufacturer	Takeda Pharmaceutical Company Limited 800.828.2088 takeda.com gammagard.com	Kedrion Biopharma, Inc. 855.353.7466 kedrion.us gammaked.com	Grifols 888.694.2686 grifols.com gamunex-c.com		
Indications	PI (adults & peds 2+ yrs.) MMN (adults, maintenance therapy) CIDP (adult)	PI (adults & peds 2+ yrs.) ITP (adults & children) CIDP (adults)	Pl (adults & peds 2+ yrs.) ITP (adults & children) CIDP (adults)		
Contraindications	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy		
Form	Liquid	Liquid	Liquid		
Concentration	10%	10%	10%		
Sizes	1/2.5/5/10/20/30 g	1/2.5/5/10/20 g	1/2.5/5/10/20/40 g		
Virus removal method	Cohn-Oncley cold ethanol fractionation, cation/ anion exchange chromatography, solvent/deter- gent, nanofiltration, low pH incubation	Cold ethanol fractionation, caprylate precipita- tion/filtration, anion exchange chromatography, low pH incubation	Cold ethanol fractionation, caprylate precipita- tion/filtration, anion exchange chromatography, low pH incubation		
IgA content	37 μg/mL	46 μg/mL	46 μg/mL		
Stabilizer	Glycine	Glycine	Glycine		
Sugar content	None	None	None		
Sodium content	None	Unspecified	Unspecified		
рН	4.6-5.1	4.0-4.5	4.0-4.5		
Osmolality (mOsmol/kg)	240-300	258	258		
Half-life (days)	31-42	35.7	35.7		
Administration	PI/MMN: IV: Begin at 0.8 mg/kg/min for first 30 minutes, if tolerated, gradually increase every 30 minutes to maximum infusion rate of 8 mg/kg/min. (MMN: May advance to 9 mg/kg/min as tolerated). PI: SubQ: Begin 1 week after last IVIG dose. (≥ 40kg) begin with 30 mL/site @ 20 mL/hr/site; maintenance 30 mL/site @ 20 mL/hr/site. (< 40kg) begin with 20 mL/site @ 15 mL/hr/ site; maintenance 20 mL/site @ 15-20 mL/hr/ site.	PI/ITP: IV: Begin at 1 mg/kg/min for first 30 minutes, if tolerated, gradually increase to maximum infusion rate of 8 mg/kg/min. CIDP: IV: Begin at 2 mg/kg/min for first 30 minutes, if tolerated, gradually increase to maximum infusion rate of 8 mg/kg/min. PI: SubQ: Begin 1 week after last IVIG dose. (adults) infuse @ 20 mL/hr/site; (children/adolescents ≥ 25kg) initially infuse @ 15 mL/hr/site; maintenance @ 20 mL/hr/site; (children/ adolescents < 25kg) infuse @ 10 mL/hr/site.	 PI/ITP: IV: Begin at 1 mg/kg/min for first 30 minutes, if tolerated, gradually increase to maximum infusion rate of 8 mg/kg/min. CIDP: IV: Begin at 2 mg/kg/min for first 30 minutes, if tolerated, gradually increase to maximum infusion rate of 8 mg/kg/min. PI: SubQ: Begin 1 week after last IVIG dose. (adults) infuse @ 20 mL/hr/site; (children/adolescents ≥ 25kg) initially infuse @ 15 mL/hr/site; maintenance @ 20 mL/hr/site; (children/adolescents < 25kg) infuse @ 10 mL/hr/site. 		
Shelf-life/storage requirements	Refrigerated 2 to 8°C [36 to 46°F] for 36 months OR room temperature up to 25°C [77°F] for 24 months	Refrigerated 2 to 8°C [36 to 46°F] for 36 months OR room temperature up to 25°C [77°F] for 6 months	Refrigerated 2 to 8°C [36 to 46°F] for 36 months OR room temperature up to 25°C [77°F] for 6 months		

Multiple brands of IG therapies for personalized care

	Intravenous							
Product	Panzyga°	Gammaplex®	Octagam®	Privigen®	Bivigam®	Asceniv*	Alyglo [*]	Gammagard [®] S/D Low IgA
Manufacturer	Octapharma USA, Inc. 866.766.4860 octapharmausa.com panzyga.info	Bio Products Laboratory Limited 866.398.0825 bpl.us.com gammaplex.com	Octapharma USA, Inc. 866.766.4860 octapharmausa.com octagamus.net	CSL Behring AG 800.504.5434 cslbehring.com privigen.com	ADMA Biologics 800.458.4244 admabiologics.com bivigam.com	ADMA Biologics 800.458.4244 admabiologics.com asceniv.com	GC Biopharma 833.426.6426 gcbiopharma.us alyglo.com	Takeda Pharmaceutical Company Limited 800.828.2088 takeda.com
Indications	PI (adults & peds 2+ yrs.) Chronic ITP (adults)	5% - PI (adults & peds 2+ yrs.); Chronic ITP 10% - PI (adults); Chronic ITP (adults)	5% - Pl 10% - Chronic ITP (adults)	Pl Chronic ITP (15+ yrs.) CIDP (adults)	Indicated for the treatment of adults and pediatric patients 2+ yrs with primary humoral immunodeficiency	Indicated for the treatment of primary humoral immunodefi- ciency (PI) in adults and adolescents (12 to 17 years of age)	Indicated for the treatment of primary humoral immuno- deficiency (PI) in adults	PI CLL Chronic ITP Kawasaki syndrome
Contraindications	IgA deficiency with antibodies to IgA & history of sensitivity; history of anaphylactic or severe systemic reactions to IG therapy	IgA deficiency with antibodies to IgA & history of sensitivity; hereditary intoler- ance to fructose; history of anaphylactic or severe systemic reactions to IG therapy	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy; hypersensitivity to corn (5%)	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy; hyperprolinemia		History of anaphylactic or severe systemic reactions to human immunoglobulin; IgA-de- ficient patients with antibodies to IgA and a history of hypersensitivity	History of anaphylactic or severe system- ic reactions to human immunoglobulin; gA-deficient patients with antibodies against IgA or a history of hypersensitivity	IgA deficiency with antibodies to IgA; history of anaphylactic or severe sys- temic reactions to IG therapy
Form	Liquid	Liquid	Liquid	Liquid	Liquid	Liquid	Liquid	Lyophilized
Concentration	10%	5/10%	5/10%	10%	10%	10%	10%	5/10%
Sizes	1/2.5/5/10/20/30 g	5/10/20 g	5%: 1/2.5/5/10/25 g 10%: 2/5/10/20/30 g	5/10/20/40 g	5/10 g	5 g	5/10/20 g	2.5/5/10 g
Virus removal method	Cold ethanol fractionation, solvent/de- tergent, ion exchange chromatography, nanofiltration	Solvent/detergent, nanofiltration, low pH incubation, cold ethanol fractionation, ion exchange chromatography	Cold ethanol fractionation, solvent/detergent, pH incubation, ultrafiltration, ion exchange chromatography	Cold ethanol fractionation, octanoic acid fractionation, anion exchange chromatog- raphy, immunoaffinity chromatography	Cold ethanol fractionation, solvent/deter- gent treatment, and 35nm virus filtration.	Cold ethanol fractionation, solvent/detergent treatment, and 35nm virus filtration.	Fractionation I+III, solvent/detergent treatment, and Nanofiltration	Cold ethanol fractionation, ultrafiltration, ion exchange chromatography, solvent/ detergent
IgA content	100 µg/mL	5%: <10 μg/mL 10%: < 20 μg/mL	5%: ≤ 200 µg/mL 10%: 106 µg/mL	≤ 25 µg/mL	≤ 200 µg/mL of IgA	≤ 200 µg/mL of IgA	≤100 mcg/mL of IgA	<1 µg/mL
Stabilizer	Glycine	5%: D-sorbitol, glycine, polysorbate 80 10%: glycine, polysorbate 80	Maltose	L-proline	Glycine	Glycine	Glycine	Albumin (human), glycine, glucose, PEG, tri-n-butyl phosphate, octoxynol, polysor- bate 80
Sugar content	None	5%: D-sorbitol (no sucrose) 10%: None	Maltose (no sucrose)	None	None	None	None	20 mg/mL glucose
Sodium content	Trace	5%: 30-50 mmol/L 10%: < 30mmol/L	≤30 mmol/L	Trace	0.100-0.140 M sodium chloride	0.100-0.140 M sodium chloride	Unspecified	5%: approx 8.5 mg/mL
рН	4.5-5.0	5%: 4.8-5.1 10%: 4.9-5.2	5%: 5.1-6.0 10%: 4.5-5.0	4.6-5.0	4.0-4.6	4.0-4.6	4.5-5.5	6.8±0.4
Osmolality (mOsmol/kg)	240-310	5%: 420-500 10%: 280	310-380	240-440	370-510	370-510	Unspecified	Unspecified
Half-life (days)	26-39	41-42	40.7	27.6-45.4	30	39.7 ± 11.6	29.6 ± 11	37.7±15
Administration	minutes; if tolerated, gradually increase every 15-30 minutes to maximum infusion rate of 8 mg/kg/min. New patients or ITP treatment, up to a	 5% infusion rate: IV: Begin at 0.5 mg/kg/min for first 15 minutes; if tolerated, gradually increase every 15 minutes to maximum infusion rate of 4 mg/kg/min. 10% infusion rate: IV: Begin at 0.5 mg/kg/min for first 15 minutes; if tolerated, gradually increase every 15 minutes to maximum infusion rate of 8 mg/kg/min. 	 5% infusion rate: IV: Begin at 0.5 mg/kg/min for first 30 minutes; if tolerated, gradually increase to maximum infusion rate of 3.33 mg/kg/min. 10% infusion rate: IV: Begin at 1 mg/kg/min for first 30 minutes; if tolerated, gradually increase to maximum infusion rate of 12 mg/kg/min. 	IV: Begin at 0.5 mg/kg/min; if tolerated, gradually increase to maximum infusion	IV: Begin at 0.5 mg/kg/min (0.005 mL/ kg/min) for first 10 min, Increase every 20 minutes (if tolerated) by 0.8 mg/kg/min up to 6 mg/kg/min	Ⅳ: Begin at 0.5 mg/kg/min (0.005 mL/kg/min) for the first 15 minutes, Increase gradually every 15 minutes (if toler- ated) up to 8 mg/kg/min (0.08 mL/kg/min)	N: 1st Infusion: Begin at 1 mg/kg/min; Double the infusion rate every 30 minutes (if tolerated) up to 8 mg/kg/min (0.08 mL/kg/min) 2nd Infusion: Begin at 2 mg/kg/min; Double the infusion rate every 15 min- utes (if tolerated) up to 8 mg/kg/min (0.08 mL/kg/min)	 5% infusion rate: IV: Begin at 0.5 mL/kg/hr, if tolerated, gradually increase to maximum infusion rate of 4 mL/kg/hr. 10% infusion rate: IV: Patient who tolerates the 5% concentration can infuse 10% solution; begin at 0.5 mL/kg/hr, if tolerated gradually increase to maximum infusion rate of 8 mL/kg/hr.
Shelf-life/storage requirements	Refrigerated 2 to 8°C [36 to 46°F] for 24 months from date of manufacture; within shelf-life, may store at $\leq 25^{\circ}$ C [77°F] for up to 9 months. After storage at $\leq 25^{\circ}$ C [77°F], either use immediately or discard	5/10%: Room temperature at 2 to 25°C [36 to 77°F] for up to 36 months	5%: Room temperature at 2 to 25°C [36 to 77°F] for up to 24 months 10%: Refrigerated 2 to 8°C [36 to 46°F] for 24 months from date of manufacture; room temperature up to 9 months at \leq 25°C [77°F] within first 12 months of shelf-life	36 months	Store at 2° to 8°C (36° to 46°F) for up to 36 months from date of manufacture. Within the first 24 months of shelf-life, product may be stored up to 4 weeks at $\leq 25^{\circ}$ C (77°F). After 24 months, product may be stored at $\leq 25^{\circ}$ C up to 2 weeks, until expiry. After storage at room temperature product must be used or discarded.	Store at 2°-8°C (36°-46°F) for up to 36 months from date of manufacture. Within the first 24 months of shelf-life, product may be stored up to 4 weeks at $\leq 25^{\circ}$ C (77° F). After 24 months, product may be stored at $\leq 25^{\circ}$ C up to 2 weeks, until expiry. After storage at room tempera- ture, product must be used or discarded.	Store ALYGLO in the refrigerator or at room temperature. Refrigeration: 2°C to 8°C [36°F to 46°F] for up to 36 months. Room Temperature: 8°C to 25°C [46°F to 77°F] for up to 24 months.	Room temperature; not to exceed 25°C [77°F]

Multiple brands of IG therapies for personalized care

	Subcutaneous				
Product	Cutaquig®	Hizentra®	Xembify®	Cuvitru®	HyQvia®
Manufacturer	Octapharma USA, Inc. 866.766.4860 octapharmausa.com cutaquigus.com	CSL Behring AG 800.504.5434 cslbehring.com hizentra.com	Grifols 888.694.2686 grifols.com xembify.com	Takeda Pharmaceutical Company Limited 800.828.2088 takeda.com cuvitru.com	Takeda Pharmaceutical Company Limited 800.828.2088 takeda.com hyqvia.com
Indications	PI (adults)	PI (adults & peds 2+ yrs.) CIDP (adults; maintenance therapy)	PI (adults & peds 2+ yrs.)	PI (adults & peds 2+ yrs.)	Pl (adults & peds 2+ yrs.) CIDP (adults)
Contraindications	IgA deficiency with antibodies to IgA & history of sensitivity; history of anaphylactic or severe systemic reactions to IG therapy	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy or components of Hizentra; hyperprolinemia	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy or inactive ingredients such as polysorbate 80	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy	IgA deficiency with antibodies to IgA; history of anaphylactic or severe systemic reactions to IG therapy; known systemic hypersensitivity to hyaluronidase and albumin (human)
Form	Liquid	Liquid	Liquid	Liquid	Liquid
Concentration	16.5%	20%	20%	20%	10%
Sizes	1/1.65/2/3.3/4/8 g	1/2/4/10 g	1/2/4/10 g	1/2/4/8/10 g	2.5/5/10/20/30 g
Virus removal method	Cold ethanol fractionation, solvent/ detergent, pH incubation, ultrafiltration, ion exchange chromatography	Cold ethanol fractionation, octanoic acid frac- tionation, anion exchange chromatography, pH incubation, nanofiltration	Cold ethanol fractionation, caprylate precipita- tion/filtration, anion exchange chromatography, low pH incubation	Cohn-Oncley cold fractionation, cation/anion exchange chromatography, solvent/detergent, nanofiltration, low pH incubation	Cohn-Oncley cold ethanol fractionation, cation/ anion exchange chromatography, solvent/deter- gent, nanofiltration, low pH incubation
IgA content	≤600 µg/mL	≤50 µg/mL	≤70 μg/mL	80 μg/mL	37 μg/mL
Stabilizer	Maltose	L-proline, polysorbate 80	Glycine, polysorbate 80	Glycine	Glycine
Sugar content	Maltose (no sucrose)	None	None	None	None
Sodium content	≤30 mmol/L	Trace	Trace	None	Recombinant human hyaluronidase compo- nent: 8.5 mg/mL sodium chloride 1.78 mg/mL sodium phosphate 0.17 mg/mL sodium hydroxide IG component: None
рН	5-5.5	4.6-5.2	4.1 to 4.8	4.6-5.1	Recombinant human hyaluronidase component: 7.4 IG component: 4.6-5.1
Osmolality (mOsmol/kg)	310-380	210-290	280-404	280-292	Recombinant human hyaluronidase component: 290-350 IG component: 240-300
Half-life (days)	Unspecified	Unspecified	Unspecified	Unspecified	50-68.6
Administration	PI: SubQ: Begin 1 week after last IVIG dose; first 6 infusions @ ≤25 mL/site @ ≤20 mL/ hr/site; maintenance ≤40 mL/site @ ≤25 mL/hr/site.	PI: SubQ: Begin 1-2 weeks after last IVIG dose, depending on dosing frequency. Initial 15 mL/site @ 15 mL/hr/site; maintenance 25 mL/site @ 25 mL/hr/site. CIDP: SubQ: Begin 1-2 weeks after last IVIG dose, depending on dosing frequency. Initial 20 mL/site @ 20 mL/hr/site; maintenance 50 mL/site @ 50 mL/hr/site.	PI: SubQ: Begin 1 week after last IVIG dose; ≤25 mL/site @ ≤25 mL/hr/site; max sites ≤6; site distance administered ≥2 inches apart.	PI: SubQ: Begin 1 week after last IVIG dose; (≥40kg) first 2 infusions @ ≤60 mL/site @ 10-20 mL/hr/site; maintenance ≤60 mL/site @ ≤60 mL/ hr/site; (<40kg) first 2 infusions @ ≤20 mL/site @ 10-20 mL/hr/site; maintenance ≤60 mL/site @ ≤ 60 mL/hr/site.	PI: SubQ: Begin 1 week after last IVIG dose; administer recombinant human hyaluronidase at initial rate/site @ 1-2 mL/min as tolerated. (≥40kg) administer up to 600 mL/site @ rates as shown in table 3 of package insert; (<40kg) administer up to 300 mL/site @ rates as shown in table 3 of package insert.
Shelf-life/ storage requirements	Refrigerated 2 to 8°C [36 to 46°F] for 24 months from date of manufacture OR room temperature up to 25°C [77°F] for 6 months during its shelf-life	Room temperature up to 25°C [77°F] for 30 months	Refrigerated 2 to 8°C [36 to 46°F] OR room temperature up to 25°C [77°F] for 6 months	Room temperature up to 25°C [77°F] for 12 months OR refrigerated 2 to 8°C [36 to 46°F] for up to 36 months	Refrigerated 2 to 8°C [36 to 46°F] for 36 months OR room temperature up to 25°C [77°F] for 3 months during the first 24 months from date of manufacture



I have a team of people who have embraced my journey, and will take that journey with me.

- Kara, Option Care Health CVID patient





See how Kara, regained her independence and joy with the help of Option Care Health's infusion therapy services. Scan the QR code to watch her inspiring story.

Contact Option Care Health to Refer a Patient!

Phone: 877.974.4844 Fax: 877.974.4845

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option care health®

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