Impact of virtual training upon patients with chronic inflammatory demyelinating polyneuropathy and primary immunodeficiency receiving subcutaneous immunoglobulin self-administration training during the coronavirus disease (COVID-19) pandemic



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Introduction

Chronic inflammatory demyelinating polyneuropathy (CIDP)

- CIDP is a progressive immune-mediated disorder of the peripheral nerves attributed to demyelination and impaired signal conduction in motor and/or sensory nerves1
- Clinical features include symmetric distal sensory and motor disorders that have proximal and distal weakness
- If left untreated, CIDP can limit patients' activity and quality of life

Primary immunodeficiencies (PIs)

- PIs comprise more than 350 different types of inherited diseases, characterized by impaired or absent components of the immune system²
- · Clinical features include increased rate and severity of infections, immune dysregulation with autoimmune disease, and malignancy³

Treating CIDP and PIs with subcutaneous immunoglobulin (SCIg)

- SCIg is approved for maintenance therapy in patients with CIDP (adults only) or PIs (adults and pediatric patients)⁴⁻⁶
- Compared with intravenous immunoglobulin (IVIg), the benefits of SCIg include self-administration in settings convenient for the patient, flexible dosing, and lower reported systemic reactions⁷⁻⁹
- Patients are often trained to self-administer by specialty pharmacy nurses in their own homes; the success of this training can affect whether the patient remains on SCIg or reverts back to their previous treatment⁹
- During the coronavirus disease 2019 (COVID-19) pandemic, an increasing number of patients transitioning to SCIg received virtual self-administration training

Objective

To assess the impact and feasibility of virtual SCIg self-administration training upon patients with CIDP or PIs during the COVID-19 pandemic

Methods

- · This was a retrospective study utilizing data collected by the Specialty Pharmacy Nurse Network (SPNN) between October 2019 and December 2020
- Patients were identified via International Classification of Diseases (ICD)-10 codes G61.81 (CIDP) or D80.0-84.9 (PIs)
- Training mode for individual patients was identified as either live, virtual, or a combination of live and virtual sessions ('combined')
- · For the purpose of this analysis, 'improved infusion parameters' was defined as an increase in either volume or rate per site, or a decrease in the number of infusion sites required between a patient's first and final session
- Data collected from (up to five) nurse-supervised training sessions were recorded:
- Number and mode of training sessions; age, gender, and BMI; training success and discontinuation rates; reasons for discontinuation; and infusion parameters from each session
- Patients who discontinued SCIg training stopped receiving SCIg therapy, and reverted to IVIg treatment; patients who completed SCIg training no longer required self-administration training by nurses

Results

Cohort characteristics

- A total of 573 patients attending 1–5 SCIg training sessions were analyzed
- The cohort was comprised 120 (21%) and 453 (79%) patients with CIDP and Pls, respectively
- The majority of training (84%) was conducted live at patients' homes (**Table 1**)
- The remaining patients were trained either fully virtually (8%) or via a combination (8%) (Table 1)
- Patients who trained virtually were slightly younger compared with those who trained live or via a combination (Table 1)
- In total, 542 (95%) patients successfully completed SCIg training
- Training success and discontinuation rates were generally comparable irrespective of training mode (**Figure 1**)
- Thirty-one (5%) SCIg discontinuations occurred during the observation period (29 trained live and two trained virtually, **Figure 1**)
- Reasons for discontinuation included: adverse events (n=12), unable to infuse independently (n=9), patient/doctor decision (n=7), worsening condition (n=1), hospitalization (n=1), or insurance issues (n=1)

Table 1: Patient characteristics by training mode Virtual, Combined All patients, Characteristic Category n=484 n=46 n=43 N=573 120 (21) 101 (21) 11 (24) 8 (19) 453 (79) 383 (79 35 (76) 35 (81) 48.1 ± 21.1 48.0 ± 21.4 45.8 ± 19.2 52.1 ± 19.3 Age (years) 357 (64) 299 (63) 32 (70) 26 (65) Female Gender, n (%) Male 201 (36) 173 (37) 14 (30) 14 (35) BMI (kg/m²) 28.5 ± 8.0 27.8 ± 7.2 27.9 ± 6.2

All values are mean + SD unless otherwise stated.

Combined training mode was defined as a combination of live and virtual training sessions.

BMI, body mass index; CIDP, chronic inflammatory demyelinating polyneuropathy; Pls, primary immunodeficiencies,

Figure 1: Impact of training mode on SCIg training success

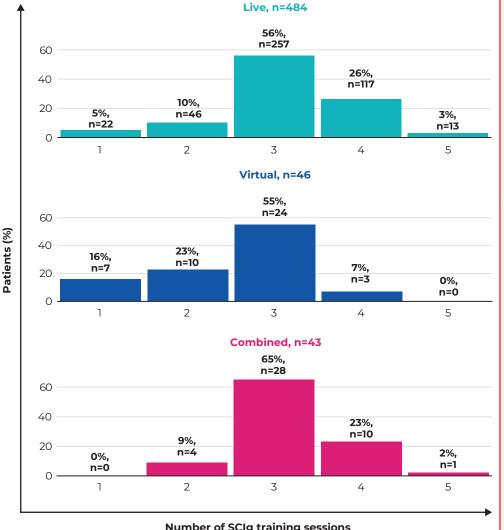


Combined training sessions were defined as a combination of live and virtual training sessions. Patients who discontinued SCIg training stopped receiving SCIg therapy and reverted to IVIg treatment; patients who completed SCIg training no longer required self-administration training by nurses.

SCIg training requirements

- Overall, most patients required three training sessions irrespective of training mode (**Figure 2**)
- Of the patients who successfully completed training, 29 (5%) required one training session, 60 (11%) required two training sessions, 309 (57%) required three training sessions, 130 (24%) required four training sessions, and 14 (3%) required five training sessions
- Patients who trained virtually were significantly more likely to require ≤3 sessions compared with patients who trained live or combined (93% vs. 71% and 74%, respectively; p=0.006)

Figure 2: Impact of the training mode on the number of sessions required by patients to complete SCIg training and become independent users of SCIg



Number of SCIg training sessions

Combined training sessions were defined as a combination of live and virtual training sessions. Due to rounding, values may not total 100%. SCIq, subcutaneous immunoglobulin

SCIg infusion parameters following training



The majority of successfully trained patients improved their infusion parameters between their first and final training session irrespective of the training mode (81%, n=440)



On average, patients who successfully completed training increased their infusion volume per site by 45% (mean ± standard deviation [SD], 6±7 mL/site) and their infusion rate per site by 46% (5±5 mL/hr/site) between their first and final training session

No differences were observed between training modes

Discussion

- Overall, patients who trained virtually were slightly younger and required fewer training sessions, compared with those who trained live or in combination
- Training success, discontinuation rates, and improvements in infusion parameters were largely comparable irrespective of training mode
- Nurse and patient confidence and experience with virtual vs. live SCIq training may have an impact upon a patient's ability to learn self-administration more quickly and with fewer discontinuations
- As such, nurse training techniques may need to differ depending on
- A limitation of this study was that fewer patients received virtual (or combined) training, compared with the number of patients who trained live

Conclusions

- Virtual (or combined) training can offer a flexible training option and appears to be as effective as live training in patients with CIDP or PIs
- More data are needed to further explore the association between training modes, ease of learning, and nursing techniques used

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Acknowledgements

The analysis, interpretation and writing of the poster for this study was funded by CSL Behring. Editorial assistance was provided by Meridian HealthComms funded by CSL Behring. We thank Cheri Jackson (Specialty Pharmacy Nursing Network [SPNN]) for her valuable insights and contributions to the study.



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