



A.S.P.E.N.'s National Patient
Registry for Nutrition Care

Essential nutrition goes into these patients through HPN.

Now, critical information will come out.

Sustain offers an unprecedented opportunity for your organization to help us collect valuable data about consumers who are on HPN. The information entered into the registry will ultimately help your organization make better-informed patient treatment decisions, support the appropriate use of nutrition support therapy, and help translate results into improved patient outcomes.

Introducing Sustain™ LLC, A.S.P.E.N.'s new online patient registry for individuals receiving nutrition support therapy. **Sustain** is the innovative tool that nutrition-support providers have sought: it allows clinicians to enter patient data, review and analyze results, benchmark outcomes against the national aggregate, post research questions and publish clinical research based on registry findings.

In phase one, **Sustain** will collect data on patients receiving Home Parenteral Nutrition (HPN). Subsequent phases will allow expansion to other populations and therapies.



Don, 70



Gail, 71



Raphael, 6



American Society for Parenteral
and Enteral Nutrition

Find out how you can participate →
www.nutritioncare.org/sustain • 800-727-456



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Sustain needs information about your patients who are on HPN



Mike, and Andrew, 16



Cyndi, and Ronnie, 61



Mallory, 25 and Maisy, 17

Currently in the United States, little is known about the annual utilization of HPN. **Sustain** will help generate data about therapy utilization, outcomes, and patient demographics which will help all consumers, past, present, and future.

Why participate?

- Contribute to prospective outcomes data development in the field of parenteral nutrition
- Help increase awareness of parenteral nutrition in the public realm
- Provide data that offers useful benchmarks and outcomes research
- Help improve clinical practice with the application of best practices found in the multi-site contributions of patient experiences
- Receive periodic reports on your patients as compared to other sites enrolled in order to benchmark your data and clinical care.

Participant benefits

- Benchmark patient data against the national aggregate
- Post research questions
- Review and analyze results
- Publish clinical research based on registry findings

Who can participate?

Any clinician who discharges HPN patients from the hospital or who cares for them in the home is eligible. Individuals must successfully complete the enrollment process which includes Institutional Review Board (IRB) approval and signing of a Participant Agreement. After data is submitted, each site will be allowed to analyze and publish their own series and benchmark against the total pooled data.

**For more information visit
www.nutritioncare.org/sustain
or contact Peggi Guenter at
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