

# The Site Council

## Healthy Relationships with Healthy Sites

Over 1,100 clinical research sites, including over 50 important site networks, AMCs and health systems, agree that the clinical research enterprise can succeed only when study sponsors and CROs have healthy relationships with healthy sites. To that end, we have created the Site Council Bill of Rights to clarify site priorities for sponsors and CROs. Sites must earn these rights by conducting clinical studies in a safe, ethical, efficient, high-quality and timely manner.

We invite clinical research sites and site networks to join the Site Council. There are no membership dues. The only member time commitment – which is modest – is briefly talking about the Site Council Bill of Rights with study sponsors and CROs. When study sponsors and CROs consistently hear from important sites about the Site Council Bill of Rights, they will have to take it seriously.

The Site Council Bill of Rights includes six fundamental rights:

- The Right to Essential Information
- The Right to Professional Autonomy
- The Right to Fair & Timely Compensation
- The Right to Proficient Governance
- The Right to Patient Centricity
- The Right to a Collaborative Relationship

These fundamental rights are supported by 24 examples that have been serious problems for over 40 years. Let's try a new, collaborative approach to solving them. The patients can't wait.

**To learn more, visit [www.sitecouncil.org](http://www.sitecouncil.org) today.**