

Topics

1. Biomarkers
2. NLP
3. Clinical Metrics System
4. Rose Score
5. Rose Clinical Growth Playbook buildout

2021 RPM rules changes

(after Public Health Emergency (PHE) regulations are lifted)

1. The mHealth technology supplied to a patient in an RPM program **must be defined as a medical device by FDA.**
2. Data **must be electronically (i.e., automatically) collected and transmitted** rather than self-reported.
3. After the PHE ends, **16 days of data must be collected** and transmitted every 30 days.
4. **Only physicians and NPPs who are eligible to furnish E/M services** may bill RPM services.
5. “interactive communication” for purposes of RPM requires, **at a minimum, a real-time synchronous, two-way audio interaction** that is capable of being enhanced with video or other kinds of data transmission.

