

One Committee to Review Them All: A Single, Multi-Disciplinary

COVID-19 Research Committee

Jocelyn Migliacci, MA, Sara Hanley, MSW, and Ann Rodavitch, MA
Memorial Sloan Kettering Cancer Center

Background

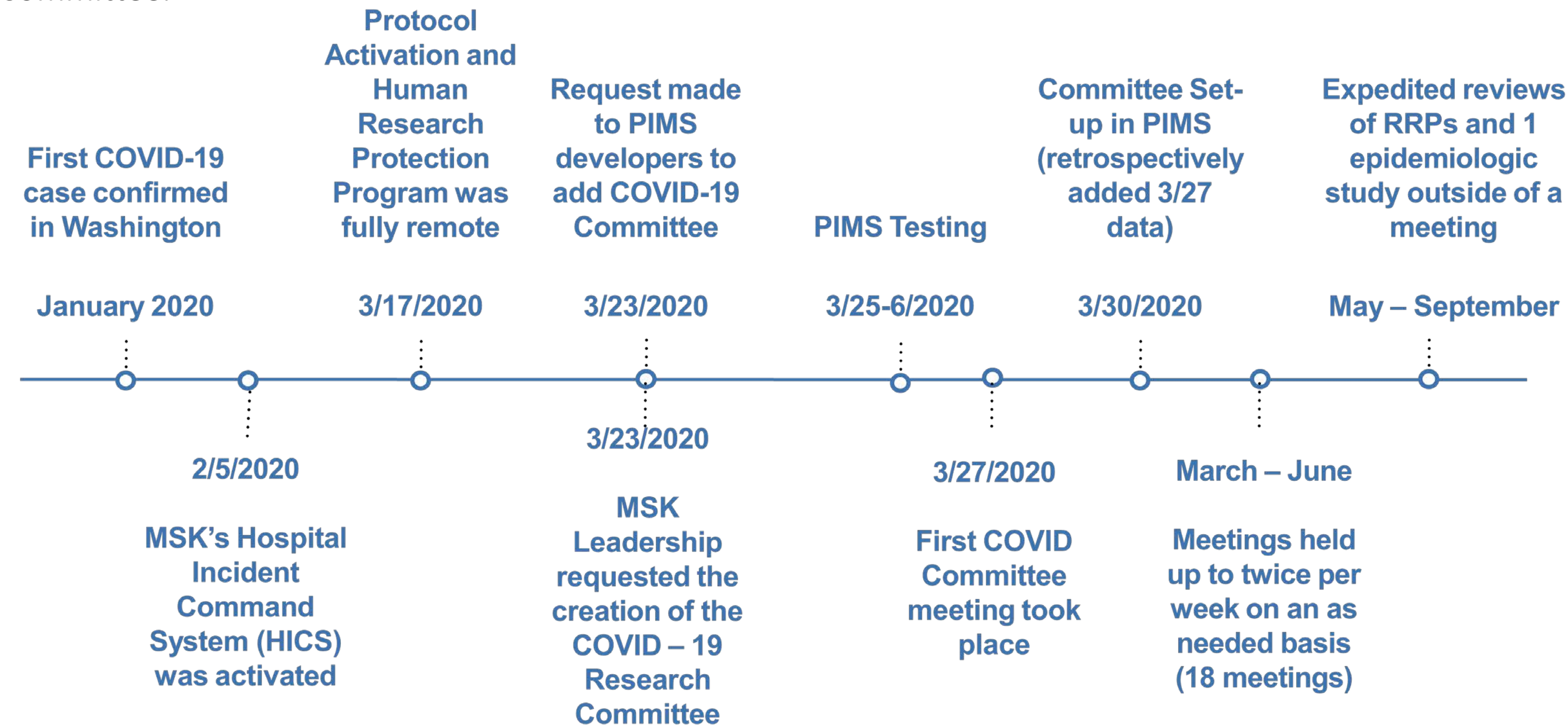
On 1/20/2020 when the first case of a novel coronavirus (COVID-19) was confirmed in Washington, its major impact was unknown. MSK's Hospital Incident Command System (HICS) was activated on 2/5, with our first COVID-19 case identified in early March. By 3/17, our Protocol Activation and Human Research Protection Program was fully remote and on 3/23, MSK leadership requested the creation of the COVID-19 Research Committee.

Goal

The goal was to create a COVID-19 Committee as a **"one-stop" committee**, providing comprehensive review of clinical research related to COVID-19, including scientific review mandated by the Cancer Center Support Grant (CCSG) guidelines, prior to IRB review.

Creation of the COVID-19 Committee

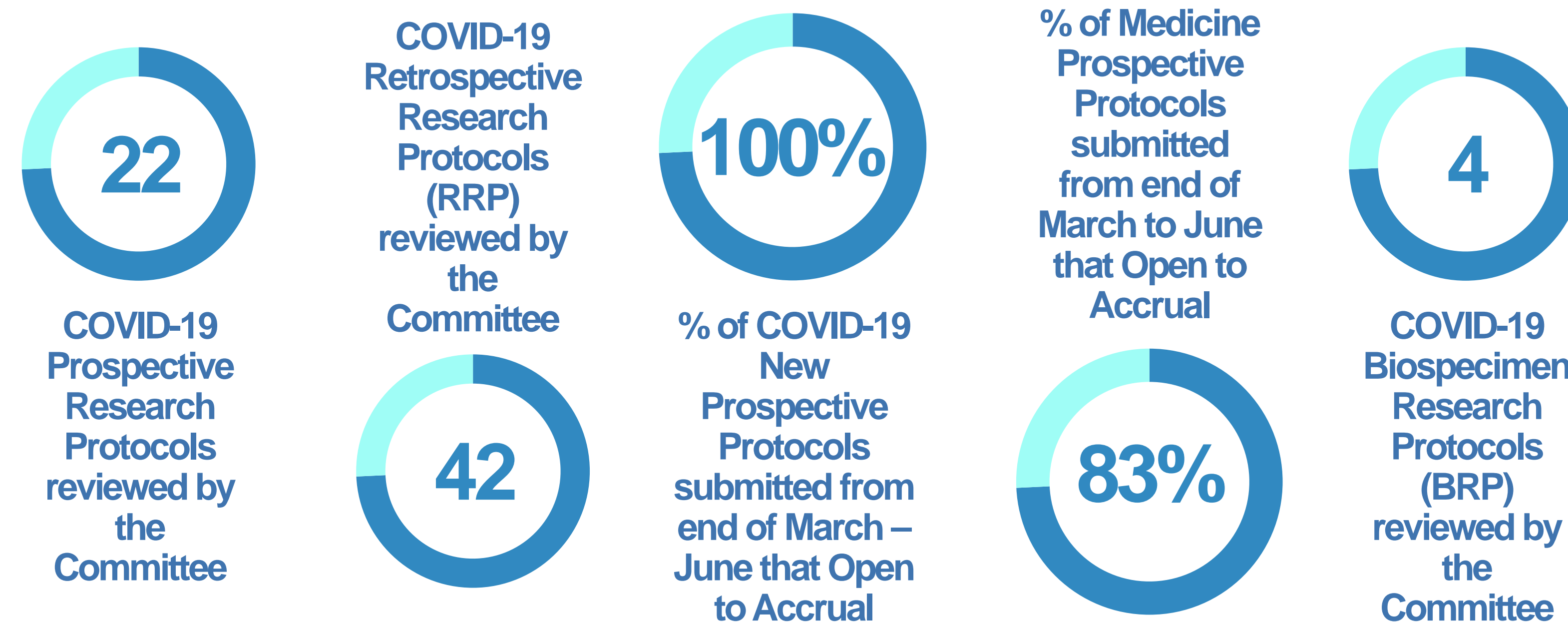
The figure below illustrates the quick timeline of events leading up to and after the creation of the COVID-19 Committee.



MSK's Protocol Information Management System (PIMS) was leveraged to efficiently manage, and track COVID-19 research. The COVID-19 Committee was created within **7 days**. Typically, it would take several months to roll out this type of large enhancement.

- Of the 8 therapeutic protocols that opened to accrual, 5 protocols have enrolled **146 participants** (in total)
- The first patient on each of the 5 studies was enrolled within **≤ 1 day** of each study opening to accrual.

Metrics (updated as of June 2021)



Role of the Committee

- ❖ To prioritize and expedite all clinical research related to COVID-19 in support of the institutional effort to rapidly activate therapeutic and other COVID-19 related research.
- ❖ To monitor the COVID-19 research portfolio to prevent overlap of efforts.
- ❖ To list all studies on the Clinical Research Portal for transparency.
- ❖ To bring together faculty from multiple disciplines, disease management groups and departments, including Physicians, Statisticians, Nurses, Pharmacists, Legal, IRB Leadership, etc.

Impressive decrease in Time to Activation (TTA) and Time to IRB Approval (TTIA) for the protocols reviewed by the COVID-19 Committee

COMMITTEE	Median TTA	Median TTIA
COVID-19 COMMITTEE	44 days	27 days
MEDICINE COMMITTEE	194 days	119 days

Next Steps

- Our unit is exploring how to utilize multidisciplinary committees to reduce the time to activation for all research protocols.
- One option we are exploring is to group protocols by disease and create a group of committees by Disease Management Team.