



Singapore Healthcare  
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# Safe and Organised Clinical Trial Specific Requisites & Medication Procurement, Storage and Utilisation through 6S & SmPC (Summary of Product Consumables)



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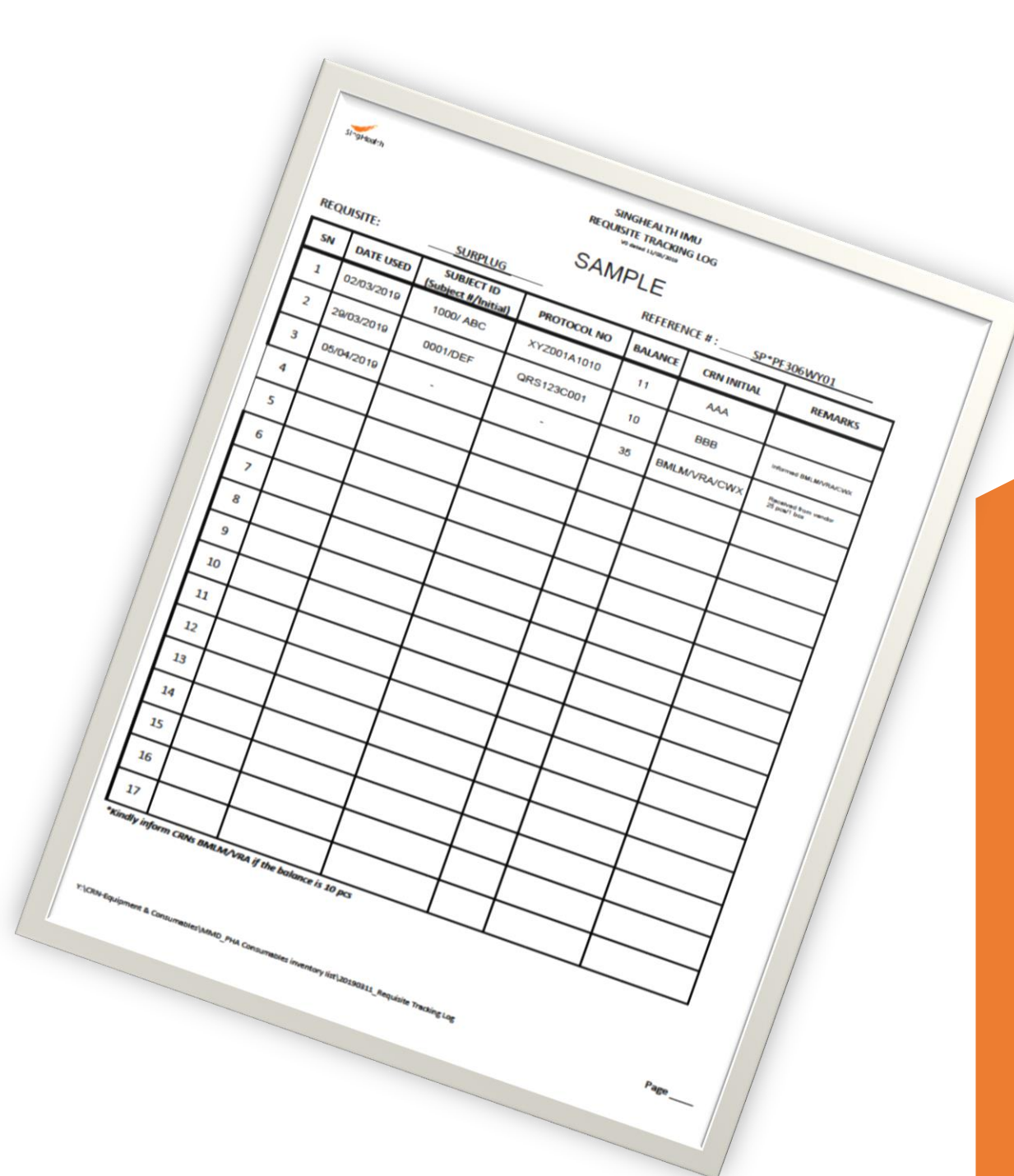
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## AIM

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To establish the importance of 6S & SmPC to ensure compliance to processes, forward-thinking procurement and trial preparation for multi-disciplinary clinical trials, leading to subject safety and trial success in IMU.



## METHODOLOGY

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**SmPC**  
Vital specifications and compatibility of requisites to Investigational Product (IP) were collated, for clear communication and smoother trial preparation.

### 6S

**Sort**  
**Set in Order**  
**Sustain**

**Safety**  
**Standardise**  
**Shine**



## INTRODUCTION

1

Clear communication is vital for subject safety in a clinical research setting. 6S & SmPC were adapted to achieve efficient execution of protocol specific procedures for smoother workflow & trial success.



6S

SmPC



## IMPACT

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- ✓ ORGANISED Trial - specific requisites storage
- ✓ EFFECTIVE internal / external communication process
- ✓ STANDARDISED guidelines and workflow
- ✓ REDUCED wastages
- ✓ MAINTAINED Subject SAFETY
- ✓ TIME EFFICIENT
- ✓ COST EFFECTIVE



## CONCLUSION

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- Expedited trial preparation
- Familiarisation with physical layouts
- Timelines met as planned
- Achieved trial success
- Cost-effective – supply and demand were met without lapses



## RESULTS

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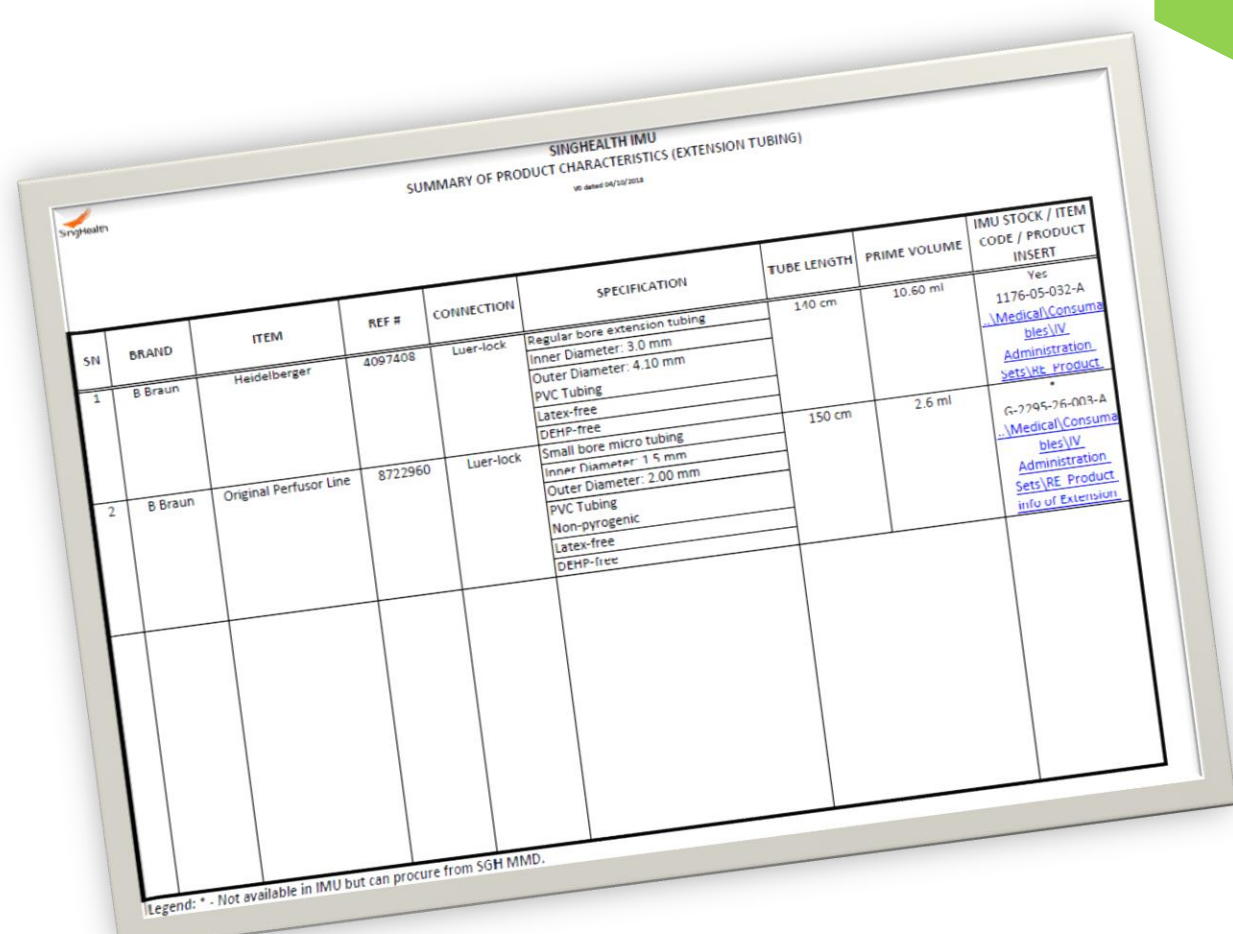
- Storage of items was strategically organized based on frequency of usage
- Par levels set for efficient tracking and procurement
- Cost effective and procurement timeline achieved



BEFORE



AFTER



## REFERENCE

Bradbury, J. (2019) 6S Safety implementation. Retrieved from <https://www.graphicproducts.com/articles/6s-safety-implementation/>