

Pilot Physiotherapy Exercise Program for Chemotherapy-Induced Peripheral Neuropathy (CIPN) in Patients with Nasopharyngeal Carcinoma

HA Convention 2026 | Oral Presentation



Dr. Ken LEE, Advanced Practice Physiotherapist



AH – Physiotherapy, Tuen Mun Hospital, NTWC



Background: Why CIPN Matters?

Most Prevalent neurological complication of chemotherapy (50–90% affected)

NPC Patients (Gemcitabine-Cisplatin + CRT): Historical incidence up to **81.7%** (Zhang et al., 2025)

Symmetric "Stocking & Glove" distribution: Affects sensory, motor & autonomic nerves

Long-term Impact: 30% experience symptoms ≥ 1 year post-treatment

Clinical Consequence: Dose reductions, delays, or discontinuation of chemotherapy leading to compromising outcomes

Limited Treatment: Duloxetine is the ONLY ASCO-recommended agent

Solution: Non-pharmacological exercise intervention is a promising strategy

CIPN Distribution Pattern



● Affected Areas

Clinical Context at Tuen Mun Hospital

Setting: Integrated Chemotherapy and Day Care Centre serving locoregionally advanced NPC patients

Treatment Regimen: Gemcitabine + Cisplatin induction chemotherapy → Concurrent Chemoradiotherapy (CRT)

Growing Burden: Increasing cancer survivorship leads to higher prevalence of long-term CIPN complications

Identified Gap: No structured physiotherapy program existed for CIPN prevention or management

Opportunity: Physiotherapy can address CIPN through tailored exercise and patient empowerment



Target Population

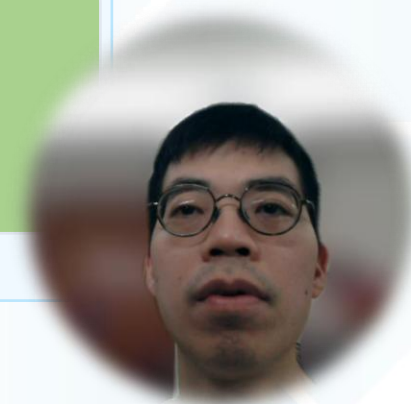
Locoregionally Advanced NPC Patients

The Challenge

Unmet need for non-pharmacological CIPN management

The Solution

Structured Physiotherapy Exercise Program



Study Objectives

1. Feasibility

Evaluate the feasibility of a tailored physiotherapy exercise program for CIPN in NPC patients.

2. Preliminary Efficacy

- Reducing CIPN incidence and severity
- Maintaining physical function (strength & tolerance)
- Preserving quality of life (QOL)
- Preventing chemotherapy dose modifications

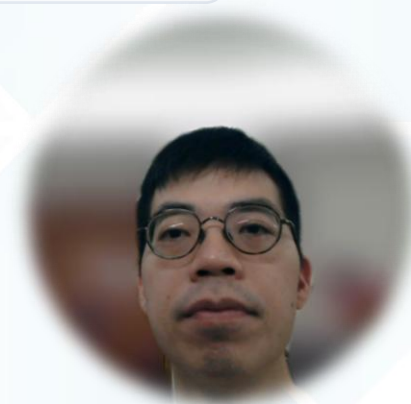
3. Patient Satisfaction

Evaluate acceptance of the hybrid delivery model (face-to-face + telehealth).

Primary Goal



TARGETING BETTER
OUTCOMES



Methodology: Study Design & Participants

Design: Prospective pilot study

Diagnosis: Locoregionally advanced Nasopharyngeal Carcinoma (NPC)

Treatment Regimen:

Gemcitabine-Cisplatin induction chemotherapy

Concurrent Chemoradiotherapy (CRT)

Inclusion Criteria:

Age 18–75 years

Histologically confirmed NPC

No prior peripheral neuropathy

Cohort Overview

 **N = 22**
Total Enrolled

 **19 M / 3 F**
Gender Distribution

 **52.2 ± 10.4**
Mean Age (Years)

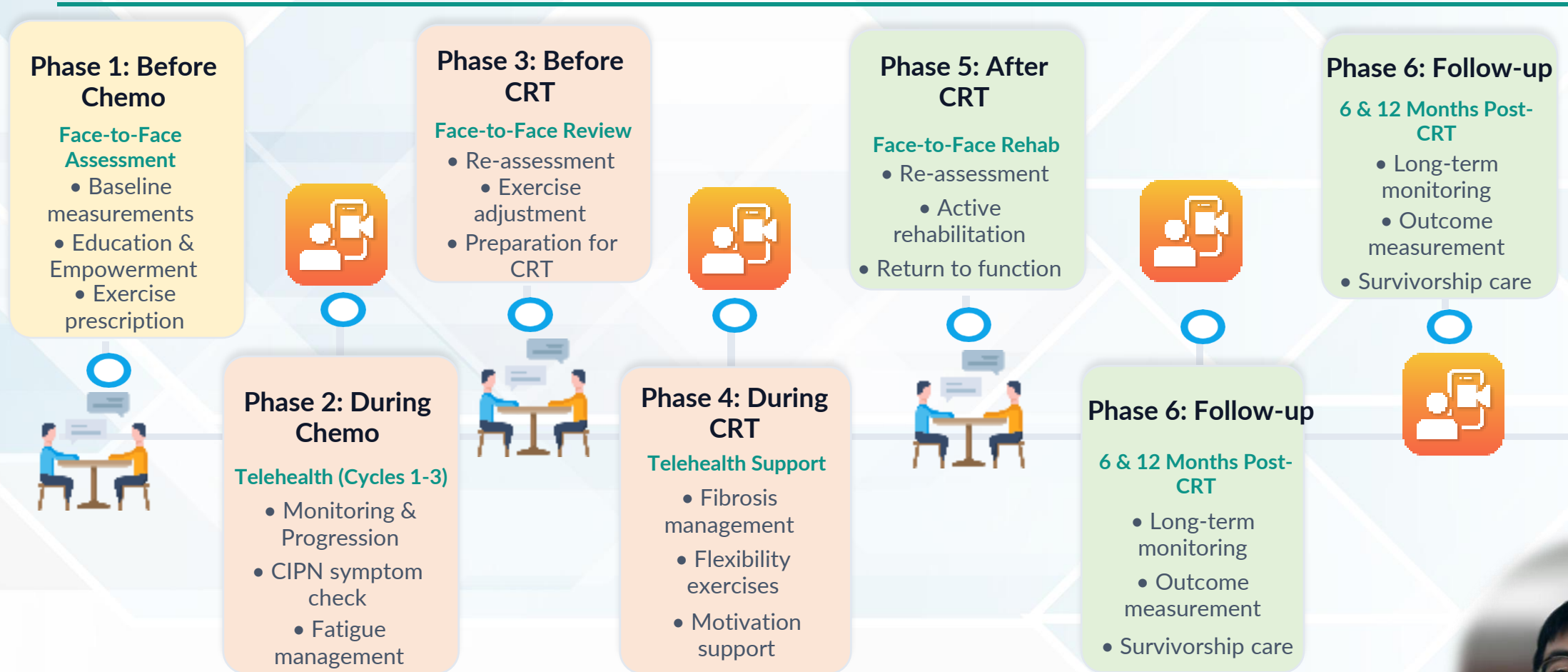
Completion Rate



Completion Rate: 12 of 22 patients (54.5%) completed full post-treatment assessment



Methodology: 6-Phase Patient Journey (7 Sessions)



Patient Empowerment Package

- Pedometer
- Two point discriminator
- Logbook
- Theraband
- Pamphlet



Methodology: Tailored Exercise Prescription



Aerobic Exercise

- **Modality:** Walking or Brisk Walking
- **Intensity:** 60–85% Heart Rate Reserve (HRR)
- **Adjustment:** Intensity modulated based on chemotherapy fatigue cycles



Pedometer



Two point discriminator



Elastic Band



Exercise log book



Balance Training

- **Progression:** Static → Dynamic balance tasks
- **Target:** Addresses CIPN-related proprioceptive loss & fall risk
- **Safety:** Performed near stable support surface



Flexibility & Nerve Gliding

- **Neurodynamics:** Specific nerve gliding exercises to maintain nerve mobility
- **Management:** Stretching for post-radiation fibrosis (neck/shoulder)



Reciprocal Pulley



Exercise Pamphlet



Resistance Training

- **Tool:** Elastic band
- **Dose:** 1–2 key exercises, 3–4 days/week
- **Progression:** Increased resistance as strength improves



Methodology: Outcome Measures

Clinical Outcomes



CIPN Incidence & Grade Primary
CTCAE v5.0 Grading Scale



Chemotherapy Delivery
Dose reductions, delays, or discontinuation



Nerve Conduction (NCS/EMG)
Objective assessment of nerve function



Exercise Tolerance
6-Minute Walk Test (6MWT)



Sensory Function
Crude Touch Monofilament Testing



Grip Strength
Handheld Dynamometry



Lower Limb Strength
30-Second Chair Stand Test

Patient-Reported Outcomes



Neurotoxicity & QOL
FACT/GOG-Ntx Scale (Chinese version)



Patient Satisfaction
Numeric Rating Scale (0-10)

Assessment Schedule

Assessments
conducted at Baseline
(Pre-Chemo), Post-
CRT 3 month



Results: Impressive Clinical Outcomes

CIPN Incidence

50%

ALL Grade 1
Study Group
(n=12)

↓ Markedly Reduced Severity

81.7%

Historical Control
(Zhang et al., 2025)

Chemo Delivery

0

Dose Reductions
due to CIPN



3 reductions occurred
for other reasons

NCS / EMG Findings (n=12)

91.7%

Developed Sensory-
Predominant Axonopathy

● 41.7% below normal limits

Clinical Interpretation

Despite objective evidence of subclinical nerve injury (axonopathy), patients remained **asymptomatic or mild (Grade 1)**.

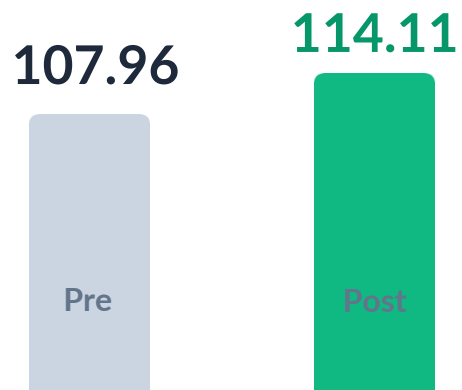
Conclusion: Exercise may mitigate SYMPTOMATIC manifestation even when nerve injury occurs.



Results: QOL & Functional Outcomes

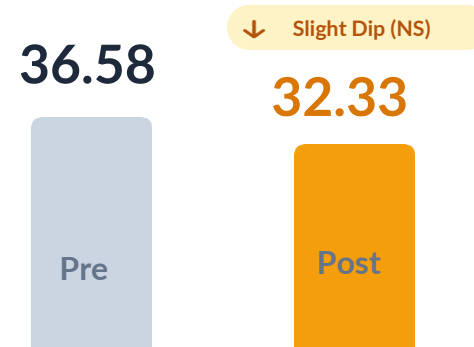
Overall Quality of Life

FACT/GOG-Ntx Total Score



Neurotoxicity Subscale

Higher score = Better outcome



Physical Function

Physical capacity remained **STABLE** throughout treatment period.



Leg Strength
Stable



Walk Distance
Stable

★ Patient Satisfaction



Hybrid Delivery Model
Face-to-Face + Telehealth

9.25



Discussion & Implications

Clinical Mechanism: Early exercise mitigates **symptomatic CIPN** even when subclinical nerve injury (NCS) occurs.

Oncological Benefit: Zero treatment delays due to CIPN supports **uninterrupted cancer treatment** for better outcomes.

QOL Value: Improvement in Quality of Life despite active chemotherapy demonstrates the program's protective effect.

Service Model: The **Hybrid Model** (Face-to-Face + Telehealth) is feasible, highly accepted, and resource-efficient.

Scalability: Protocol is adaptable to other HA cancer centers, addressing the growing survivorship demand.

Clinical Efficacy

Decoupling nerve injury from symptoms allows patients to maintain function.

System Efficiency

Telehealth integration reduces hospital visits while maintaining close monitoring.

Wide Applicability

Physiotherapy-led model enhances the oncology Multi-Disciplinary Team (MDT).



Conclusion & Future Directions

Conclusion

- ✓ The pilot CIPN physiotherapy exercise program is **FEASIBLE and SAFE** for NPC patients.
- ✓ **EFFECTIVE** in reducing symptomatic CIPN incidence **50% vs 81.7%**
- ✓ **SUCCESSFUL** in preserving physical function and Quality of Life.
- ✓ Highly **ACCEPTED** by patients (satisfaction 9.25/10) with hybrid delivery model.
- ✓ Supports **uninterrupted chemotherapy delivery** (zero delays due to CIPN).

Future Directions

- 1 Expand Scope: Include patients with diverse cancer types and agents linked to CIPN (e.g., Oxaliplatin, Paclitaxel).
- 2 Enhance Training: Implement efficient patient education models, such as chemotherapy-focused classes.
- 3 Standardize Protocols: Establish a unified physiotherapy protocol for CIPN management.
- 4 Implement at Scale: Roll out service-wide across NTWC and extend to all HA clusters.



Acknowledgements & References

Acknowledgements

Authors & Co-investigators

Lee YT, Liu YT, Kwok PW, Yeung I, Wong CY, Mui WH, Chan LY, Ho PS, Tsoi SY, Kwan WS, Lam WS, Mak MY

Supervisor

Mr. Vincent LAM, SPT (Physio)

Department

AH – Physiotherapy, Tuen Mun Hospital, NTWC

Special Thanks

To all participating patients for their commitment and the Oncology Team at TMH for their collaboration.

Key References

- 1 Zhang et al., 2025 - Random forest model analysis of influencing factors of chemotherapy-induced peripheral neuropathy in patients with nasopharyngeal carcinoma: a cross-sectional survey study. *Frontiers in oncology* vol. 15 1548742. 18 Jul. 2025, doi:10.3389/fonc.2025.1548742
- 2 ASCO Guidelines – Loprinzi, Charles L et al. “Prevention and Management of Chemotherapy-Induced Peripheral Neuropathy in Survivors of Adult Cancers: ASCO Guideline Update.” *Journal of clinical oncology : official journal of the American Society of Clinical Oncology* vol. 38,28 (2020): 3325-3348. doi:10.1200/JCO.20.01399

