固体分光Ⅰ 擬一次元 NaV₂O₄の高分解能 ARPES

東北大院理[^],東北大 WPI^B,物材機構^c T. Qian^A,佐藤宇史[^],中山耕輔[^],荒金俊行[^], P. Richard^B,新井正男^c, 山浦一成^c,室町英治^c,高橋 隆^{A,B}

Angle-resolved photoemission study of quasi-one-dimensional metallic compound NaV₂O₄

Dept. Physics, Tohoku Univ.^A, WPI, Tohoku Univ.^B, NIMS ^C

T. Qian^A, T. Sato^A, K. Nakayama^A, T. Arakane^A, P. Richard^B, M. Arai^C,

K. Yamaura^C, E. Takayama-Muromachi^C and T. Takahashi^{A,B}

It is recently reported that a newly synthesized vanadium oxide NaV_2O_4 takes a quasi-one-dimensional (1D) metallic ground state at low temperatures [1]. Our previous photoemission measurements have revealed the power-law behavior near Fermi level (E_F) characteristic of the quasi-1D materials [2]. To clarify the quasi-1D correlation in NaV_2O_4 , it is strongly desired to clarify the momentum dependence of the electronic structure near E_F .

carried out We have an angle-resolved photoemission spectroscopy (ARPES) study of NaV₂O₄. The ARPES spectra at 20 K reveal a clear dispersive feature within 0.3 eV of $E_{\rm F}$ and a Fermi surface crossing along the chain direction with the absence of a Fermi edge. As seen in Fig. 1, the Fermi surface mapping gives an open Fermi surface with a finite curvature perpendicular to the chain direction. The Fermi surface topology not only provides direct evidence for the guasi-1D correlation in NaV_2O_4 , but also reveals the existence of finite interchain hopping, which leads to weak but finite spectral weight at $E_{\rm F}$. We also compared the experimental band structure with the local spin density approximation band calculations at the ferromagnetic and nonmagnetic states. The experimental bands can be well understood based on the moderate exchange splitting.

- [1] K. Yamaura *et al.*, Phys. Rev. Lett. **99**, 196601 (2007).
- [2] T. Qian et al., J. Phys. Soc. Jpn. (in press).



Fig. 1. Experimental Fermi surface of NaV_2O_4 measured at 20 K with 100-eV photon energy.