

学歴

- 1976年3月 東京大学理学部化学科卒業
1978年3月 東京大学大学院理学系研究科化学専攻修士課程修了
1983年6月 理学博士（東京大学）

職歴

- 1978年3月 東京大学物性研究所 表面物性部門教務職員
1983年3月 東京大学理学部物理学教室助手
1990年3月 大阪大学基礎工学部物性物理学教室助教授
1991年4月 東京大学物性研究所軌道放射物性部門客員助教授（1992年3月まで）
1994年8月 アメリカ合衆国、カリフォルニア州バークレー市ローレンスバークレー
研究所に客員研究員として滞在（1995年5月まで）
1997年4月 奈良先端科学技術大学院大学物質創成科学研究科教授（評議員、学長補佐、
研究科長など兼務）
2004年7月 立命館大学 COE 推進機構特別招聘教授
2019年3月 奈良先端科学技術大学院大学教授を停年退職
2019年4月 公益財団法人豊田理化学研究所フェロー
2019年4月 日本原子力研究開発機構客員研究員（併任）

2. これまでの研究成果

1) 昭和 51 年 4 月～昭和 53 年 3 月 東京大学理学部化学教室 朽津耕三研究室にて、修士論文研究として、気体分子 (O_2 , N_2 , CCl_4 , As_4) による低速電子散乱の実験的、理論的解析。

修士論文研究として数百 eV の電子線の気体分子からの微分散乱断面積 (散乱角度分布) の測定と解析を行った。広い角度範囲にわたって散乱電子の強度を一点一点測定した。微分散乱断面積の中には分子内原子からの散乱波の干渉パターンが現れ、それを解析して分子構造を求めることができるが、分子内多重散乱が起きると構造解析に誤差が生じる。分子内多重散乱の大きさがどの程度か、どのようなパターンとして現れるか、などを明らかにするため、多重散乱の効果が顕著に現れることが予想される数百 eV の低エネルギーにおいて、いくつかの気体分子 (O_2 , N_2 , CCl_4 , As_4) について実験的及び理論的に解析した。理論計算は、独立原子モデルに分子内多重散乱、分極、電子交換などを考慮して行った。原子内及び原子間の干渉パターンが鈍り、絶対値が下がる効果を初めて明らかにした。

2) 昭和 53 年 3 月～昭和 58 年 3 月 東京大学物性研究所表面物性部門 村田好正研究室にて、固体表面 (Si の酸化, 遷移金属シリサイド, S/Ni (001)) の放射光光電子分光・光電子回折による研究。

博士論文タイトル「Experimental and Theoretical Studies of Elastic Electron Scattering at Intermediate Energies and Photoemission Studies of Reactions on Semiconductor Surfaces Using Synchrotron Radiation」

当時は世界で初めての放射光専用リングが東大の田無キャンパスに建設され、表面研究への応用が可能になった時期であった。博士論文研究として、放射光を用いた光電子分光・光電子回折の研究を日本で初めて行った。Si は酸素励起種によって効率よく良質の酸化膜を作ること、Ni などの遷移金属シリサイドの電子状態は各成分のバンドの重ねあわせでよく理解できること、Ni (001) 上の S からの光電子回折パターンから、S の吸着位置は最表面 Ni の 1.3 Å 上であることなどを明らかにした。光電子回折は、1) の電子の干渉効果を固体表面に応用したものである。これ以降、「光電子回折・光電子ホログラフィー」による表面の原子配列構造の研究を推進してきた。

ここでも広い角度範囲にわたって光電子の強度を一点一点測定していたが、光電子では強度が弱いためこの測定は特に時間がかかり、角度分布が一度に測定できる二次元検出器の必要性を強く意識した。

3) 昭和 58 年 3 月～平成 2 年 3 月 東京大学理学部物理学教室 井野正三研究室にて、二次元表示型球面鏡分析器の発明、RHEED を用いた表面構造の研究、超高真空走査電子顕微鏡の開発

その当時は、米国で回転楕円鏡型の二次元表示型分析器が Eastman によって発明されて使用されていたが、取り出し角度範囲は $\pm 43^\circ$ 程度が限界であり、また得られる像が歪んでいるという問題があった。そこで、新しい型の二次元表示型分析器の作成を目指して電子の軌道解析を繰り返して行った。その結果、球面を利用してメッシュと電極の間隔を 2 倍程度に大きくすると厳密に収束する解が得られ、かつ像の歪みはまったく無く、取り出し立体角はいくらでも大きく取れる (表面の場合は 180°) 二次元表示型球面鏡分析器を発明することができた (図 1) [1]。この分析器を用いると、試料から放出された電子の角度分布を、広い立体角に渡って二次元的に歪み無く一度に表示できる。一方向だけ取り込む従来の分析器に比べると一万倍程度効率が上がる。この研究は、日本化学会若

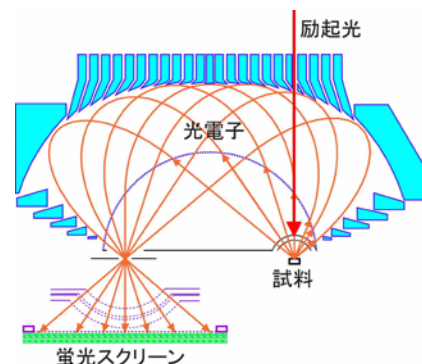


図 1 二次元表示型球面鏡分析器

い世代の特別講演に選ばれた。

この時期のその他の研究としては、井野正三教授の指導で RHEED による表面構造解析、超高真空電子顕微鏡の作成（電子顕微鏡学会賞受賞）などが挙げられる。

4) 平成 2 年 3 月～平成 9 年 4 月 大阪大学基礎工学部 物性物理学教室 菅滋正研究室にて、放射光二次元光電子分光、光電子ホログラフィーによる表面研究、円偏光光電子回折における前方散乱ピークの回転の発見

1990 年代になると放射光技術も進み、円偏光放射光が使えるようになったので、円偏光励起光電子回折の実験を企画した。円偏光励起ではスピンは変わらないが軌道角運動量が変わるので、光電子が軌道角運動量を持つことを予想した。X線を原子に照射すると、原子の中の電子は光電子となって飛び出す。その運動エネルギーが数百 eV 以上のとき、図 2 のように、光電子を放出した原子と、その隣の原子とを結ぶ方向に前方散乱ピークと呼ばれるピークが観測される。光電子が古典的な軌道角運動量を持てば、光電子回折パターンの中の前方散乱ピークが図のように回転するという予想を立てて実験を行ったところ、予想通りの回転を観測することができた。この結果は、これまで概念的にしか捉えられていなかった軌道角運動量を初めて直接観測したことになり、非光学活性物質の円 2 色性を初めて観測したことにもなったので、応用物理学会賞を戴いた[2]。この「前方散乱ピークの回転」という現象の解釈の確立のため、米国の放射光施設 ALS でも実験と解析を行った。

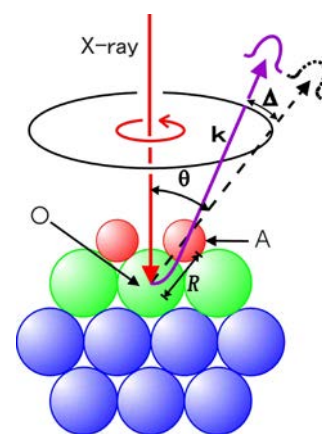


図 2 前方散乱ピークの回転

また、固体のエネルギーバンドを、放射光を使って 2 次的に測定すると、バンドを構成している電子の軌道や結合状態を明らかにできることを見出した[3]。光電子構造因子という概念を創出し、これを用いると、グラファイトでは π バンドが p_z 結合軌道から構成されていることが簡単にわかり、Bi 系超伝導体では $d_{x^2-y^2}$ 軌道がフェルミ面を作っていることを明らかにすることができた。

また、当時米国で開発された光電子ホログラフィーの研究を始めた。これは、円偏光ではない軟 X 線で励起した光電子角度分布をフーリエ変換することによって、着目原子の周りの原子配列を立体的に再現するものである。日本で初めての論文をいくつか報告した。

5) 平成 9 年 4 月～平成 31 年 3 月 奈良先端科学技術大学院大学物質創成科学研究科にて、偏光放射光二次元光電子分光による表面電子状態、立体原子写真法の発明

奈良先端科学技術大学院大学に移ってからは、文科省からの予算の他に、下記の外部資金をいただいて研究を進めてきたので、資金ごとにとめる。

- 平成 10 年度～平成 13 年度 特別推進研究「二次元光電子分光と電子物性」(研究代表者) 239,000,000 円

研究目的は、図 1 の二次元光電子分光器のエネルギー分解能を高め、固体の電子エネルギーバンドを 3 次的に測定し、物性のメカニズムを電子論的に解明しようというものであった。高エネルギー分解能の二次元表示型球面鏡分析器を作製し、それを用いて図 3 に示すグラファイトの 3 次的なバンド構造の測定、原子軌道の帰属、ウムクラップ過程の解析にも成功した。多くのベストポスター賞を受け、事後評価も A であった。

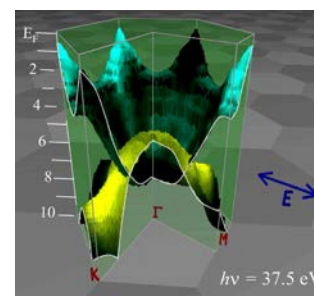


図 3 グラファイトの 3D エネルギーバンド

◎立体原子写真法の発明

この研究を進めている中で、前述の「前方散乱ピークの回転」の回転角を視差角に見立てて、着目する原子の周りの原子配列を立体的に直視できるという「**立体原子写真法**」を発明することができた[4]。解析してみると、ほとんどゆがみのない立体写真が得られることがわかった。

立体写真を測定する装置は、上記の改良型「二次元表示型球面鏡分析器」である。図4 (a), (b)は、この装置で測定した立体写真の例で、W (110) 面からのW 4 f 内殻光電子の放出角度分布パターンである。兵庫県の放射光施設 SPring-8 で使用できる円偏光 X 線を励起光として用いた。光電子の運動エネルギーは 800eV である。(a), (b)は、それぞれ回転の向きが右および左の円偏光を用いて測定したものである。このパターンは、図4 (d)のO原子から図4 (d)の上の方を見ていることになる。図4 (a), (b)においては、図4 (d)の原子A, B, Cによる前方散乱ピークが、図4 (c)のA, B, Cの位置に観測されている。しかしながら、(a), (b)におけるピーク位置は、結晶構造から予想される原子の方向(図4 (c)のA, B, Cの位置)より左右に少しずれていて、そのずれがO原子に人間がいて原子A, B, Cを両目で見たときの視差角と同じく、近くの原子ほど大きくなっている。従って、(a), (b)の図を左右の目でそれぞれ見ることにより、原子A, B, Cの立体配列を図4 (d)のように直接認識することができる。このずれは、光電子が円偏光の角運動量をもらうことによって実質的に角運動量を持つため、原子の周りに回転しながら放射されることによって生じることを明らかにした。

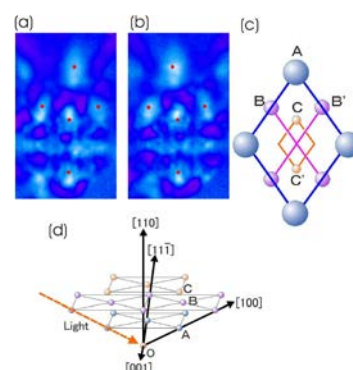


図4 W 結晶の立体写真

● 平成 14 年度～平成 19 年度 戦略的創造研究 CREST (科学技術振興機構)「ナノ構造解析のための立体原子顕微鏡の開発」(研究代表者) 235,000,000 円

この戦略的創造研究 CREST においては、上記で開発された立体原子写真法を種々の試料に応用して手法として確立すること、使いやすい測定装置を作製すること、および低倍率の機能を付加すること、の3つの目標があった。上記の立体写真を測定していた装置では、原子像が直視できる10億倍程度の高倍率の像は得られるものの、角度分布のみを測定しているため、不均一な試料のどこを見ているかを調べる低倍率の像は得られなかった。試料表面から飛び出した電子をレンズを使って焦点を結ばば低倍率の拡大像が得られるが、これまでのレンズは $\pm 2^\circ$ 程度の範囲しか結像できず、立体原子写真法に必要な $\pm 50^\circ$ のような広い角度範囲を収束することは不可能であった。そこで、非球面メッシュレンズを発明して、 $\pm 50^\circ$ に渡る角度範囲の放出電子を一点に収束できる広角対物レンズを実現し、特許を取得した。

- 平成 20 年度～平成 24 年度 基盤研究 (S) (文部科学省科学研究費補助金)「微小領域二次元光電子分光」(研究代表者) 132, 200, 000 円

この基盤研究 (S) においては、上記で発明した $\pm 50^\circ$ の広角対物レンズを用いて、低倍率の拡大像も得られて微小領域からだけの二次元光電子分光が行える高エネルギー分解能の分析器を完成することを目的とした。図 5 に組み上げた DELMA (Display-type Ellipsoidal Mesh Analyzer) を示す。図 5 の右下に示すように、試料表面の低倍率の拡大像、 $\pm 50^\circ$ の角度校正パターン、および光電子回折パターンを得ることに成功した[5]。高エネルギー分解能の測定の際には、図 5 のスクリーンを引き上げて、電子を後ろの市販の高エネルギー分解能の分析器に通す。一度に測定できる範囲は小さくなるが、試料や装置は動かさずにレンズの最後のデフレクタを走査するだけで $\pm 50^\circ$ の角度に渡ってエネルギー分解能の高い測定が可能になった。

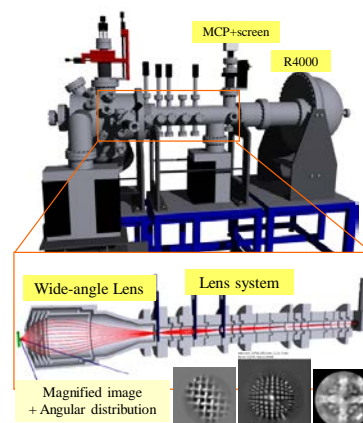


図 5 DELMA

- 平成 26 年度～平成 30 年度 新学術領域研究 (文部科学省科学研究費補助金)「3D 活性サイト科学」(領域代表者) 1, 621, 600, 000 円

前述の「光電子ホログラフィー」は、しばらくは原子像の再現精度が悪くて実用にならない状況が続いていたが、この頃までに松下らによる精度の良い解析手法が開発され、実用レベルに達していた。また、その他にも林による蛍光 X 線ホログラフィーの長足の進歩もあり、日本で原子分解能ホログラフィーの新学術領域を立ち上げる機運が高まり、平成 26 年度に採択されてスタートした。金属や半導体にドーパされた元素は、機能発現に重要な役割を果たしているが、周期構造を取らないために通常の構造解析手段である X 線回折が使えず、局所構造がわからないままで開発が試行錯誤で行われていた。この新学術では、半導体、超伝導体、たんぱく質などのバイオ物質の中で活性を担っている元素の周りの原子配列を種々の原子分解能ホログラフィーを用いて解析することを目的としている。例として、Ca を層間に入れた超伝導体グラファイトに K をドーパした試料を光電子ホログラフィーで観測した結果を図 6 に示す[6]。炭素の作るハニカム構造と、K がその 2 倍の周期で入っていることが良く分かる。他にも、Si 半導体中のヒ素ドーパントや、ヘモグロビンたんぱく質の中心で活性を担っている Fe の周りの原子配列など、多くの成果が上がっている。このように、これまで一貫して偏光の放射光を用いた二次元光電子分光を用いて、原子レベルの構造解析、および新しい解析法の開発を行ってきた。

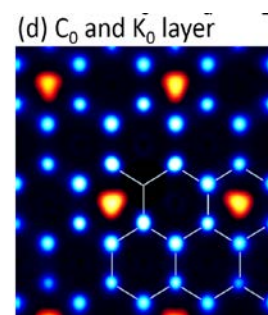


図 6 Ca と K をドーパしたグラファイト超伝導体の光電子ホログラフィー

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